

## PHENYLALANINE DERIVATIVES

5 This invention relates to a series of phenylalanine derivatives, to compositions containing them, to processes for their preparation, and to their use in medicine.

10 Over the last few years it has become increasingly clear that the physical interaction of inflammatory leukocytes with each other and other cells of the body plays an important role in regulating immune and inflammatory responses [Springer, T A. *Nature*, 346, 425, (1990); Springer, T. A. *Cell* 76, 301, (1994)]. Many of these interactions are mediated by specific cell surface molecules collectively referred to as cell adhesion molecules.

15 The adhesion molecules have been sub-divided into different groups on the basis of their structure. One family of adhesion molecules which is believed to play a particularly important role in regulating immune and inflammatory responses is the integrin family. This family of cell surface glycoproteins has a typical non-covalently linked heterodimer structure. At least 14 different integrin alpha chains and 8 different integrin beta chains  
20 have been identified [Sonnenberg, A. *Current Topics in Microbiology and Immunology*, 184, 7, (1993)]. The members of the family are typically named according to their heterodimer composition although trivial nomenclature is widespread in this field. Thus the integrin termed  $\alpha 4 \beta 1$  consists of the integrin alpha 4 chain associated with the integrin beta 1  
25 chain, but is also widely referred to as Very Late Antigen 4 or VLA4. Not all of the potential pairings of integrin alpha and beta chains have yet been observed in nature and the integrin family has been subdivided into a number of subgroups based on the pairings that have been recognised [Sonnenberg, A. *ibid*].

30 The importance of cell adhesion molecules in human leukocyte function has been further highlighted by a genetic deficiency disease called Leukocyte Adhesion Deficiency (LAD) in which one of the families of leukocyte integrins is not expressed [Marlin, S. D. *et al* *J. Exp. Med.* 164, 855 (1986)]. Patients with this disease have a reduced ability to recruit

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leukocytes to inflammatory sites and suffer recurrent infections which in extreme cases may be fatal.

5 The potential to modify adhesion molecule function in such a way as to  
beneficially modulate immune and inflammatory responses has been  
extensively investigated in animal models using specific monoclonal  
antibodies that block various functions of these molecules [e.g. Issekutz, T.  
B. J. Immunol. 3394, (1992); Li, Z. *et al* Am. J. Physiol. 263, L723, (1992);  
10 Binns, R. M. *et al* J. Immunol. 157, 4094, (1996)]. A number of  
monoclonal antibodies which block adhesion molecule function are  
currently being investigated for their therapeutic potential in human  
disease.

15 One particular integrin subgroup of interest involves the  $\alpha 4$  chain which  
can pair with two different beta chains  $\beta 1$  and  $\beta 7$  [Sonnenberg, A. *ibid*.  
The  $\alpha 4\beta 1$  pairing occurs on many circulating leukocytes (for example  
lymphocytes, monocytes and eosinophils) although it is absent or only  
present at low levels on circulating neutrophils.  $\alpha 4\beta 1$  binds to an adhesion  
molecule (Vascular Cell Adhesion Molecule-1 also known as VCAM-1)  
20 frequently up-regulated on endothelial cells at sites of inflammation  
[Osborne, L. Cell, 62, 3, (1990)]. The molecule has also been shown to  
bind to at least three sites in the matrix molecule fibronectin [Humphries,  
M. J. *et al*. Ciba Foundation Symposium, 189, 177, (1995)]. Based on  
data obtained with monoclonal antibodies in animal models it is believed  
25 that the interaction between  $\alpha 4\beta 1$  and ligands on other cells and the  
extracellular matrix plays an important role in leukocyte migration and  
activation [Yednock, T. A. *et al*, Nature, 356, 63, (1992); Podolsky, D. K.  
*et al*. J. Clin. Invest. 92, 373, (1993); Abraham, W. M. *et al*. J. Clin. Invest.  
93, 776, (1994)].

30 The integrin generated by the pairing of  $\alpha 4$  and  $\beta 7$  has been termed  
LPAM-1 [Holzmann, B and Weissman, I. EMBO J. 8, 1735, (1989)] and  
like  $\alpha 4\beta 1$ , binds to VCAM-1 and fibronectin. In addition,  $\alpha 4\beta 7$  binds to an  
adhesion molecule believed to be involved in the homing of leukocytes to  
35 mucosal tissue termed MAdCAM-1 [Berlin, C. *et al*, Cell, 74, 185, (1993)].  
The interaction between  $\alpha 4\beta 7$  and MAdCAM-1 may also be important at

sites of inflammation outside of mucosal tissue [Yang, X-D. *et al*, PNAS, 91, 12604 (1994)].

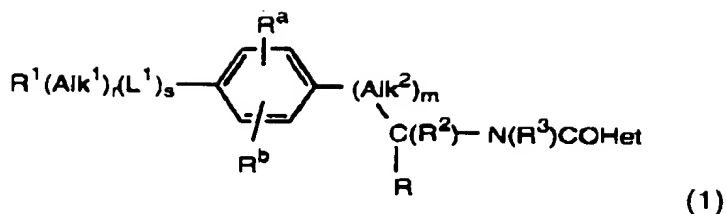
Regions of the peptide sequence recognised by  $\alpha 4\beta 1$  and  $\alpha 4\beta 7$  when they  
 5 bind to their ligands have been identified.  $\alpha 4\beta 1$  seems to recognise LDV, IDA or REDV peptide sequences in fibronectin and a QIDSP sequence in VCAM-1 [Humphries, M. J. *et al*, *ibid*] whilst  $\alpha 4\beta 7$  recognises a LDT sequence in MAdCAM-1 [Briskin, M. J. *et al*, J. Immunol. 156, 719, (1996)]. There have been several reports of inhibitors of these interactions  
 10 being designed from modifications of these short peptide sequences [Cardarelli, P. M. *et al* J. Biol. Chem. 269, 18668, (1994); Shroff, H. N. Bioorganic. Med. Chem. Lett. 6, 2495, (1996); Vanderslice, P. J. Immunol. 158, 1710, (1997)]. It has also been reported that a short peptide sequence derived from the  $\alpha 4\beta 1$  binding site in fibronectin can inhibit a  
 15 contact hypersensitivity reaction in a trinitrochlorobenzene sensitised mouse [Ferguson, T. A. *et al*, PNAS 88, 8072, (1991)].

Since the alpha 4 subgroup of integrins are predominantly expressed on  
 leukocytes their inhibition can be expected to be beneficial in a number of  
 20 immune or inflammatory disease states. However, because of the ubiquitous distribution and wide range of functions performed by other members of the integrin family it is very important to be able to identify selective inhibitors of the alpha 4 subgroup.

25 We have now found a group of compounds which are potent and selective inhibitors of  $\alpha 4$  integrins. Members of the group are able to inhibit  $\alpha 4$  integrins such as  $\alpha 4\beta 1$  and/or  $\alpha 4\beta 7$  at concentrations at which they generally have no or minimal inhibitory action on  $\alpha$  integrins of other subgroups. The compounds are thus of use in medicine, for example in  
 30 the prophylaxis and treatment of immune or inflammatory disorders as described hereinafter.

Thus according to one aspect of the invention we provide a compound of formula (1)

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wherein

R is a carboxylic acid or a derivative thereof;

- 5 R<sup>1</sup> is a hydrogen atom or a hydroxyl, straight or branched alkoxy or optionally substituted cycloaliphatic, polycycloaliphatic, heterocycloaliphatic, polyheterocycloaliphatic, aromatic or heteroaromatic group;  
 Alk<sup>1</sup> is an optionally substituted aliphatic or heteroaliphatic chain;  
 L<sup>1</sup> is a linker atom or group;

- 10 r and s, which may be the same or different, is each zero or an integer 1 provided that when r is zero R<sup>1</sup> is an optionally substituted cycloaliphatic, polycycloaliphatic, heterocycloaliphatic, polyheterocycloaliphatic, aromatic or heteroaromatic group;  
 R<sup>a</sup> and R<sup>b</sup>, which may be the same or different is each an atom or group  
 15 -L<sup>2</sup>(CH<sub>2</sub>)<sub>p</sub>L<sup>3</sup>(R<sup>c</sup>)<sub>q</sub> in which L<sup>2</sup> and L<sup>3</sup> is each a covalent bond or a linker atom or group, p is zero or the integer 1, q is an integer 1, 2 or 3 and R<sup>c</sup> is a hydrogen or halogen atom or a group selected from straight or branched alkyl, -OR<sup>d</sup> [where R<sup>d</sup> is a hydrogen atom or an optionally substituted straight or branched alkyl group], -SR<sup>d</sup>, -NR<sup>d</sup>R<sup>e</sup>, [where R<sup>e</sup> is as just defined for R<sup>d</sup> and may be the same or different], -NO<sub>2</sub>, -CN, -CO<sub>2</sub>R<sup>d</sup>,  
 20 -SO<sub>3</sub>H, -SO<sub>2</sub>R<sup>d</sup>, -OCO<sub>2</sub>R<sup>d</sup>, -CONR<sup>d</sup>R<sup>e</sup>, -OCONR<sup>d</sup>R<sup>e</sup>, -CSNR<sup>d</sup>R<sup>e</sup>, -COR<sup>d</sup>, -N(R<sup>d</sup>)COR<sup>e</sup>, N(R<sup>d</sup>)CSR<sup>e</sup>, -SO<sub>2</sub>N(R<sup>d</sup>)(R<sup>e</sup>), -N(R<sup>d</sup>)SO<sub>2</sub>R<sup>e</sup>, -N(R<sup>d</sup>)CONR<sup>e</sup>R<sup>f</sup> [where R<sup>f</sup> is a hydrogen atom or an optionally substituted straight or branched alkyl group], -N(R<sup>d</sup>)CSNR<sup>e</sup>R<sup>f</sup> or -N(R<sup>d</sup>)SO<sub>2</sub>NR<sup>e</sup>R<sup>f</sup>;

- 25 Alk<sup>2</sup> is a straight or branched alkylene chain;  
 m is zero or an integer 1;  
 R<sup>2</sup> is a hydrogen atom or a methyl group;  
 R<sup>3</sup> is a hydrogen atom or a straight or branched alkyl group;  
 Het is an optionally substituted heteroaromatic group;  
 30 and the salts, solvates, hydrates and N-oxides thereof.

It will be appreciated that compounds of formula (1) may have one or more chiral centres. Where one or more chiral centres is present, enantiomers or diastereomers may exist, and the invention is to be understood to extend to all such enantiomers, diastereomers and mixtures thereof, including racemates. Formula (1) and the formulae hereinafter are intended to represent all individual isomers and mixtures thereof, unless stated or shown otherwise.

In the compounds of formula (1), derivatives of the carboxylic acid group R include carboxylic acid esters and amides. Particular esters and amides include  $-\text{CO}_2\text{Alk}^4$  and  $-\text{CON}(\text{R}^4)_2$  groups as described herein.

When in the compounds of the invention  $\text{L}^1$  is present as a linker atom or group it may be any divalent linking atom or group. Particular examples include  $-\text{O}-$  or  $-\text{S}-$  atoms or  $-\text{C}(\text{O})-$ ,  $-\text{C}(\text{O})\text{O}-$ ,  $-\text{C}(\text{S})-$ ,  $-\text{S}(\text{O})-$ ,  $-\text{S}(\text{O})_2-$ ,  $-\text{N}(\text{R}^4)-$  [where  $\text{R}^4$  is a hydrogen atom or a straight or branched alkyl group],  $-\text{CON}(\text{R}^4)-$ ,  $-\text{OC}(\text{O})\text{N}(\text{R}^4)-$ ,  $-\text{CSN}(\text{R}^4)-$ ,  $-\text{N}(\text{R}^4)\text{CO}-$ ,  $-\text{N}(\text{R}^4)\text{C}(\text{O})\text{O}-$ ,  $-\text{N}(\text{R}^4)\text{CS}-$ ,  $-\text{S}(\text{O})\text{N}(\text{R}^4)-$ ,  $-\text{S}(\text{O})_2\text{N}(\text{R}^4)-$ ,  $-\text{N}(\text{R}^4)\text{S}(\text{O})-$ ,  $-\text{N}(\text{R}^4)\text{S}(\text{O})_2-$ ,  $-\text{N}(\text{R}^4)\text{CON}(\text{R}^4)-$ ,  $-\text{N}(\text{R}^4)\text{CSN}(\text{R}^4)-$ ,  $-\text{N}(\text{R}^4)\text{SON}(\text{R}^4)-$  or  $-\text{N}(\text{R}^4)\text{SO}_2\text{N}(\text{R}^4)-$  groups. Where the linker group contains two  $\text{R}^4$  substituents, these may be the same or different.

$\text{Alk}^2$  in the compounds of the invention may be for example a straight or branched  $\text{C}_{1-3}$ alkylene chain. Particular examples include  $-\text{CH}_2-$ ,  $-\text{CH}(\text{CH}_3)-$ ,  $-\text{C}(\text{CH}_3)_2-$  and  $-(\text{CH}_2)_2-$ .

When  $\text{R}^3$  and/or  $\text{R}^4$  in the compounds of formula (1) is a straight or branched alkyl group it may be a straight or branched  $\text{C}_{1-6}$ alkyl group, e.g. a  $\text{C}_{1-3}$ alkyl group such as a methyl or ethyl group.

When  $\text{Alk}^1$  in compounds of formula (1) is an optionally substituted aliphatic chain it may be an optionally substituted  $\text{C}_{1-10}$  aliphatic chain. Particular examples include optionally substituted straight or branched chain  $\text{C}_{1-6}$  alkyl,  $\text{C}_{2-6}$  alkenyl, or  $\text{C}_{2-6}$  alkynyl chains.

Heteroaliphatic chains represented by  $\text{Alk}^1$  include the aliphatic chains just described but with each chain additionally containing one, two, three or four heteroatoms or heteroatom-containing groups. Particular heteroatoms or groups include atoms or groups  $\text{L}^4$  where  $\text{L}^4$  is as defined above for  $\text{L}^1$  when  $\text{L}^1$  is a linker atom or group. Each  $\text{L}^4$  atom or group may interrupt the aliphatic chain, or may be positioned at its terminal carbon atom to connect the chain to the atom or group  $\text{R}^1$ .

Particular examples of aliphatic chains represented by  $\text{Alk}^1$  include optionally substituted  $-\text{CH}_2-$ ,  $-\text{CH}_2\text{CH}_2-$ ,  $-\text{CH}(\text{CH}_3)-$ ,  $-\text{C}(\text{CH}_3)_2-$ ,  $-(\text{CH}_2)_2\text{CH}_2-$ ,  $-\text{CH}(\text{CH}_3)\text{CH}_2-$ ,  $-(\text{CH}_2)_3\text{CH}_2-$ ,  $-\text{CH}(\text{CH}_3)\text{CH}_2\text{CH}_2-$ ,  $-\text{CH}_2\text{CH}(\text{CH}_3)\text{CH}_2-$ ,  $-\text{C}(\text{CH}_3)_2\text{CH}_2-$ ,  $-(\text{CH}_2)_4\text{CH}_2-$ ,  $-(\text{CH}_2)_5\text{CH}_2-$ ,  $-\text{CHCH}-$ ,  $-\text{CHCHCH}_2-$ ,  $-\text{CH}_2\text{CHCH}-$ ,  $-\text{CHCHCH}_2\text{CH}_2-$ ,  $-\text{CH}_2\text{CHCHCH}_2-$ ,  $-(\text{CH}_2)_2\text{CHCH}-$ ,  $-\text{CC}-$ ,  $-\text{CCCH}_2-$ ,  $-\text{CH}_2\text{CC}-$ ,  $-\text{CCCH}_2\text{CH}_2-$ ,  $-\text{CH}_2\text{CCCH}_2-$ , or  $-(\text{CH}_2)_2\text{CC}-$  chains. Where appropriate each of said chains may be optionally interrupted by one or two atoms and/or groups  $\text{L}^4$  to form an optionally substituted heteroaliphatic chain. Particular examples include optionally substituted  $-\text{L}^4\text{CH}_2-$ ,  $-\text{CH}_2\text{L}^4\text{CH}_2-$ ,  $-\text{L}^4(\text{CH}_2)_2-$ ,  $-\text{CH}_2\text{L}^4(\text{CH}_2)_2-$ ,  $-(\text{CH}_2)_2\text{L}^4\text{CH}_2-$ ,  $-\text{L}^4(\text{CH}_2)_3-$  and  $-(\text{CH}_2)_2\text{L}^4(\text{CH}_2)_2-$  chains. The optional substituents which may be present on aliphatic or heteroaliphatic chains represented by  $\text{Alk}^1$  include one, two, three or more substituents selected from halogen atoms, e.g. fluorine, chlorine, bromine or iodine atoms, or hydroxyl,  $\text{C}_1$ -6alkoxy, e.g. methoxy or ethoxy, thiol,  $\text{C}_1$ -6alkylthio e.g. methylthio or ethylthio, amino or substituted amino groups. Substituted amino groups include  $-\text{NHR}^4$  and  $-\text{N}(\text{R}^4)_2$  groups where  $\text{R}^4$  is a straight or branched alkyl group as defined above. Where two  $\text{R}^4$  groups are present these may be the same or different. Particular examples of substituted chains represented by  $\text{Alk}^1$  include those specific chains just described substituted by one, two, or three halogen atoms such as fluorine atoms, for example chains of the type  $-\text{CH}(\text{CF}_3)-$ ,  $-\text{C}(\text{CF}_3)_2-$ ,  $-\text{CH}_2\text{CH}(\text{CF}_3)-$ ,  $-\text{CH}_2\text{C}(\text{CF}_3)_2-$ ,  $-\text{CH}(\text{CF}_3)-$  and  $-\text{C}(\text{CF}_3)_2\text{CH}_2-$ .

Alkoxy groups represented by  $\text{R}^1$  in compounds of the invention include straight or branched  $\text{C}_1$ -6alkoxy groups such as methoxy and ethoxy groups.

When R<sup>1</sup> is present in compounds of formula (1) as an optionally substituted cycloaliphatic group it may be an optionally substituted C<sub>3-10</sub> cycloaliphatic group. Particular examples include optionally substituted C<sub>3-10</sub>cycloalkyl, e.g. C<sub>3-7</sub>cycloalkyl, C<sub>3-10</sub>cycloalkenyl e.g. C<sub>3-7</sub>cycloalkenyl or C<sub>3-10</sub>cycloalkynyl e.g. C<sub>3-7</sub>cycloalkynyl groups.

Optionally substituted heterocycloaliphatic groups represented by R<sup>1</sup> include the optionally substituted cycloaliphatic groups just described for R<sup>1</sup> but with each group additionally containing one, two, three or four heteroatoms or heteroatom-containing groups L<sup>2</sup> as just defined.

Optionally substituted polycycloaliphatic groups represented by R<sup>1</sup> include optionally substituted C<sub>7-10</sub> bi- or tricycloalkyl or C<sub>7-10</sub>bi- or tricycloalkenyl groups. Optionally substituted polyheterocycloaliphatic groups represented by R<sup>1</sup> include the optionally substituted polycycloalkyl groups just described, but with each group additionally containing one, two, three or four L<sup>2</sup> atoms or groups.

Particular examples of R<sup>1</sup> cycloaliphatic, polycycloaliphatic, heterocycloaliphatic and polyheterocycloaliphatic groups include optionally substituted cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl, cycloheptyl, 2-cyclobuten-1-yl, 2-cyclopenten-1-yl, 3-cyclopenten-1-yl, adamantyl, norbornyl, norbornenyl, pyrroline, e.g. 2- or 3-pyrrolinyl, pyrrolidinyl, pyrrolidinone, oxazolidinyl, oxazolidinone, dioxolanyl, e.g. 1,3-dioxolanyl, imidazoliny, e.g. 2-imidazoliny, imidazolidinyl, pyrazoliny, e.g. 2-pyrazoliny, pyrazolidinyl, thiazoliny, thiazolidinyl, pyranyl, e.g. 2- or 4-pyranyl, piperidinyl, piperidinone, 1,4-dioxanyl, morpholiny, morpholinone, 1,4-dithianyl, thiomorpholiny, piperazinyl, 1,3,5-trithianyl, oxaziny, e.g. 2H-1,3-, 6H-1,3-, 6H-1,2-, 2H-1,2- or 4H-1,4- oxaziny, isoxaziny, oxathiaziny, e.g. 1,2,5 or 1,2,6-oxathiaziny, or 1,3,5-oxadiaziny groups.

The optional substituents which may be present on the R<sup>1</sup> cycloaliphatic, polycycloaliphatic, heterocycloaliphatic or polyheterocycloaliphatic groups include one, two, three or more substituents represented by R<sup>5</sup> in which R<sup>5</sup> is selected from halogen atoms, e.g. fluorine, chlorine, bromine or iodine atoms, or C<sub>1-6</sub>alkyl, e.g. methyl or ethyl, haloC<sub>1-6</sub>alkyl, e.g.

halomethyl or haloethyl such as difluoromethyl or trifluoromethyl, hydroxyl, C<sub>1-6</sub>alkoxy, e.g. methoxy or ethoxy, haloC<sub>1-6</sub>alkoxy, e.g. halomethoxy or haloethoxy such as difluoromethoxy or trifluoromethoxy, thiol, C<sub>1-6</sub>alkylthio e.g. methylthio or ethylthio, -N(R<sup>4</sup>)<sub>2</sub>, -CN, -CO<sub>2</sub>R<sup>4</sup>, -NO<sub>2</sub>, -CON(R<sup>4</sup>)<sub>2</sub>,  
 5 -CSN(R<sup>4</sup>)<sub>2</sub>, -COR<sup>4</sup>, -CSN(R<sup>4</sup>)<sub>2</sub>, -N(R<sup>4</sup>)COR<sup>4</sup>, -N(R<sup>4</sup>)CSR<sup>4</sup>, -SO<sub>2</sub>N(R<sup>4</sup>)<sub>2</sub>,  
 -N(R<sup>4</sup>)SO<sub>2</sub>R<sup>4</sup>, -N(R<sup>4</sup>)CON(R<sup>4</sup>)<sub>2</sub>, -N(R<sup>4</sup>)CSN(R<sup>4</sup>) and -N(R<sup>4</sup>)SO<sub>2</sub>N(R<sup>4</sup>)<sub>2</sub> groups. In these substituents the group R<sup>4</sup> when present is a hydrogen atom or a straight or branched alkyl group as defined above. Where more than one R<sup>4</sup> group is present in a substituent each group may be the same  
 10 or different. The substituent may be present on any available carbon atom or where appropriate any nitrogen atom, in the R<sup>1</sup> group

In the compounds of formula (1), optionally substituted aromatic groups represented by the group R<sup>1</sup> include for example monocyclic or bicyclic  
 15 fused ring C<sub>6-12</sub> aromatic groups, such as phenyl, 1- or 2-naphthyl, 1- or 2-tetrahydronaphthyl, indanyl or indenyl groups, optionally substituted by one, two, three or more -L<sup>2</sup>(CH<sub>2</sub>)<sub>p</sub>L<sup>3</sup>(R<sup>c</sup>)<sub>q</sub> atoms or groups, where L<sup>2</sup>, L<sup>3</sup>, p and q are as previously defined and R<sup>c</sup> is as previously defined but is other than a hydrogen atom when L<sup>2</sup> and L<sup>3</sup> is each a covalent bond and p  
 20 is zero.

Optionally substituted heteroaromatic groups, represented by the group R<sup>1</sup> or Het in compounds of formula (1) include for example optionally substituted C<sub>1-9</sub> heteroaromatic groups containing for example one, two,  
 25 three or four heteroatoms selected from oxygen, sulphur or nitrogen atoms. In general, the heteroaromatic groups may be for example monocyclic or bicyclic fused ring heteroaromatic groups. Monocyclic heteroaromatic groups include for example five- or six-membered heteroaromatic groups containing one, two, three or four heteroatoms  
 30 selected from oxygen, sulphur or nitrogen atoms. Bicyclic heteroaromatic groups include for example nine- to thirteen-membered fused-ring heteroaromatic groups containing one, two or more heteroatoms selected from oxygen, sulphur or nitrogen atoms.

35 Particular examples of heteroaromatic groups of these types include optionally substituted pyrrolyl, furyl, thienyl, imidazolyl, N-C<sub>1-6</sub>imidazolyl,



oxazolyl, isoxazolyl, thiazolyl, isothiazolyl, pyrazolyl, 1,2,3-triazolyl, 1,2,4-triazolyl, 1,2,3-oxadiazolyl, 1,2,4-oxadiazolyl, 1,2,5-oxadiazolyl, 1,3,4-oxadiazolyl, 1,3,4-thiadiazole, pyridyl, pyrimidinyl, pyridazinyl, pyrazinyl, 1,3,5-triazinyl, 1,2,4-triazinyl, 1,2,3-triazinyl, benzofuryl, [2,3-dihydro]-  
 5 benzofuryl, benzothienyl, benzotriazolyl, indolyl, isoindolyl, benzimidazolyl, imidazo[1,2-a]pyridyl, benzothiazolyl, benzoxazolyl, benzopyranyl, [3,4-dihydro]benzopyranyl, quinazolinyl, naphthyridinyl, pyrido[3,4-b]pyridyl, pyrido[3,2-b]pyridyl, pyrido[4,3-b]pyridyl, quinolinyl, isoquinolinyl, tetrazolyl, 5,6,7,8-tetrahydroquinolinyl, 5,6,7,8-tetrahydroisoquinolinyl, and imidyl,  
 10 e.g. succinimidyl, phthalimidyl, or naphthalimidyl such as 1,8-naphthalimidyl.

Optional substituents which may be present on R<sup>1</sup> heteroaromatic groups include one, two, three or more -L<sup>2</sup>(CH<sub>2</sub>)<sub>p</sub>L<sup>3</sup>(R<sup>c</sup>)<sub>q</sub> atoms on groups as just  
 15 defined.

Examples of the substituents represented by R<sup>a</sup> and R<sup>b</sup> in compounds of formula (1) and which may be present on aromatic or heteroaromatic groups represented by R<sup>1</sup> include atoms or groups -L<sup>2</sup>(CH<sub>2</sub>)<sub>p</sub>LR<sup>c</sup>,  
 20 -L<sup>2</sup>(CH<sub>2</sub>)<sub>p</sub>R<sup>c</sup>, -L<sup>2</sup>R<sup>c</sup>, -(CH<sub>2</sub>)<sub>p</sub>R<sup>c</sup> and -R<sup>c</sup> wherein L<sup>2</sup>, (CH<sub>2</sub>)<sub>p</sub>, L and R<sup>c</sup> are as defined above. Particular examples of such substituents include -L<sup>2</sup>CH<sub>2</sub>L<sup>2</sup>R<sup>c</sup>, -L<sup>2</sup>CH(CH<sub>3</sub>)L<sup>3</sup>R<sup>c</sup>, -L<sup>2</sup>(CH<sub>2</sub>)<sub>2</sub>L<sup>3</sup>R<sup>c</sup>, -2CH<sub>2</sub>R<sup>c</sup>, -L<sup>2</sup>CH(CH<sub>3</sub>)R<sup>c</sup>, -L<sup>2</sup>(CH<sub>2</sub>)<sub>2</sub>R<sup>c</sup>, -CH<sub>2</sub>R<sup>c</sup>, -CH(CH<sub>3</sub>)R<sup>c</sup> and -(CH<sub>2</sub>)<sub>2</sub>R<sup>c</sup> groups.

Thus each of R<sup>a</sup> and R<sup>b</sup> and, where present, substituents on R<sup>1</sup> aromatic or heteroaromatic groups in compounds of the invention may be for example selected from a hydrogen atom, a halogen atom, e.g. a fluorine, chlorine, bromine or iodine atom, or a C<sub>1-6</sub>alkyl, e.g. methyl, ethyl, n-propyl, i-propyl, n-butyl or t-butyl, C<sub>1-6</sub>alkylamino, e.g. methylamino or  
 25 ethylamino, C<sub>1-6</sub>hydroxyalkyl, e.g. hydroxymethyl, hydroxyethyl or  
 30 -C(OH)(CF<sub>3</sub>)<sub>2</sub>, carboxyC<sub>1-6</sub>alkyl, e.g. carboxyethyl, C<sub>1-6</sub>alkylthio e.g. methylthio or ethylthio, carboxyC<sub>1-6</sub>alkylthio, e.g. carboxymethylthio, 2-carboxyethylthio or 3-carboxypropylthio, C<sub>1-6</sub>alkoxy, e.g. methoxy or ethoxy, hydroxyC<sub>1-6</sub>alkoxy, e.g. 2-hydroxyethoxy, haloC<sub>1-6</sub>alkyl, e.g. -CF<sub>3</sub>,  
 35 -CHF<sub>2</sub>, CH<sub>2</sub>F, haloC<sub>1-6</sub>alkoxy, e.g. -OCF<sub>3</sub>, -OCHF<sub>2</sub>, -OCH<sub>2</sub>F, C<sub>1-6</sub>alkylamino, e.g. methylamino or ethylamino, amino (-NH<sub>2</sub>), aminoC<sub>1-6</sub>alkyl, e.g.

- aminomethyl or aminoethyl, C<sub>1-6</sub>dialkylamino, e.g. dimethylamino or diethylamino, C<sub>1-6</sub>alkylaminoC<sub>1-6</sub>alkyl, e.g. ethylaminoethyl, C<sub>1-6</sub>dialkylaminoC<sub>1-6</sub>alkyl, e.g. diethylaminoethyl, aminoC<sub>1-6</sub>alkoxy, e.g. aminoethoxy, C<sub>1-6</sub>alkylaminoC<sub>1-6</sub>alkoxy, e.g. methylaminoethoxy, C<sub>1-6</sub>dialkylaminoC<sub>1-6</sub>alkoxy, e.g. dimethylaminoethoxy, diethylaminoethoxy, isopropylaminoethoxy, or dimethylaminopropoxy, nitro, cyano, amidino, hydroxyl (-OH), formyl [HC(O)-], carboxyl (-CO<sub>2</sub>H), -CO<sub>2</sub>R<sup>12</sup>, C<sub>1-6</sub>alkanoyl e.g. acetyl, thiol (-SH), thioC<sub>1-6</sub>alkyl, e.g. thiomethyl or thioethyl, sulphonyl (-SO<sub>3</sub>H), C<sub>1-6</sub>alkylsulphonyl, e.g. methylsulphonyl, aminosulphonyl (-SO<sub>2</sub>NH<sub>2</sub>), C<sub>1-6</sub>alkylaminosulphonyl, e.g. methylaminosulphonyl or ethylaminosulphonyl, C<sub>1-6</sub>dialkylaminosulphonyl, e.g. dimethylaminosulphonyl or diethylaminosulphonyl, phenylaminosulphonyl, carboxamido (-CONH<sub>2</sub>), C<sub>1-6</sub>alkylaminocarbonyl, e.g. methylaminocarbonyl or ethylaminocarbonyl, C<sub>1-6</sub>dialkylaminocarbonyl, e.g. dimethylaminocarbonyl or diethylaminocarbonyl, aminoC<sub>1-6</sub>alkylaminocarbonyl, e.g. aminoethylaminocarbonyl, C<sub>1-6</sub>dialkylaminoC<sub>1-6</sub>alkylaminocarbonyl, e.g. diethylaminoethylaminocarbonyl, aminocarbonylamino, C<sub>1-6</sub>alkylaminocarbonylamino, e.g. methylaminocarbonylamino or ethylaminocarbonylamino, C<sub>1-6</sub>dialkylaminocarbonylamino, e.g. dimethylaminocarbonylamino or diethylaminocarbonylamino, C<sub>1-6</sub>alkylaminocarbonylC<sub>1-6</sub>alkylamino, e.g. methylaminocarbonylmethylamino, aminothiocarbonylamino, C<sub>1-6</sub>alkylaminothiocarbonylamino, e.g. methylaminothiocarbonylamino or ethylaminothiocarbonylamino, C<sub>1-6</sub>dialkylaminothiocarbonylamino, e.g. dimethylaminothiocarbonylamino or diethylaminothiocarbonylamino, C<sub>1-6</sub>alkylaminothiocarbonylC<sub>1-6</sub>alkylamino, e.g. ethylaminothiocarbonylmethylamino, C<sub>1-6</sub>alkylsulphonylamino, e.g. methylsulphonylamino or ethylsulphonylamino, C<sub>1-6</sub>dialkylsulphonylamino, e.g. dimethylsulphonylamino or diethylsulphonylamino, aminosulphonylamino (-NHSO<sub>2</sub>NH<sub>2</sub>), C<sub>1-6</sub>alkylaminosulphonylamino, e.g. methylaminosulphonylamino or ethylaminosulphonylamino, C<sub>1-6</sub>dialkylaminosulphonylamino, e.g. dimethylaminosulphonylamino or diethylaminosulphonylamino, C<sub>1-6</sub>alkanoylamino, e.g. acetylamino, aminoC<sub>1-6</sub>alkanoylamino e.g. aminoacetylamino, C<sub>1-6</sub>dialkylaminoC<sub>1-6</sub>alkanoylamino, e.g. dimethylaminoacetylamino, C<sub>1-6</sub>alkanoylaminoC<sub>1-6</sub>alkyl, e.g. acetylaminomethyl, C<sub>1-6</sub>alkanoylaminoC<sub>1-6</sub>alkylamino, e.g. acetamidoethylamino, C<sub>1-</sub>

alkoxycarbonylamino, e.g. methoxycarbonylamino, ethoxycarbonylamino or t-butoxycarbonylamino group.

- Optional substituents present on the heteroaromatic groups represented by Het include one, two, three or more substituents, each selected from an atom or group  $R^6$  in which  $R^6$  is  $-R^{6a}$  or  $-\text{Alk}^3(R^{6a})_m$ , where  $R^{6a}$  is a halogen atom, or an amino ( $-\text{NH}_2$ ), substituted amino, nitro, cyano, amidino, hydroxyl ( $-\text{OH}$ ), substituted hydroxyl, formyl, carboxyl ( $-\text{CO}_2\text{H}$ ), esterified carboxyl, thiol ( $-\text{SH}$ ), substituted thiol,  $-\text{COR}^7$  [where  $R^7$  is an  $-\text{Alk}^3(R^{6a})_m$ , aryl or heteroaryl group],  $-\text{CSR}^7$ ,  $-\text{SO}_3\text{H}$ ,  $-\text{SO}_2\text{R}^7$ ,  $-\text{SO}_2\text{NH}_2$ ,  $-\text{SO}_2\text{NHR}^7$ ,  $-\text{SO}_2\text{N}(R^7)_2$ ,  $-\text{CONH}_2$ ,  $-\text{CSNH}_2$ ,  $-\text{CONHR}^7$ ,  $-\text{CSNHR}^7$ ,  $-\text{CON}(R^7)_2$ ,  $-\text{CSN}(R^7)_2$ ,  $-\text{N}(R^4)\text{SO}_2\text{R}^7$ ,  $-\text{N}(\text{SO}_2\text{R}^7)_2$ ,  $-\text{NH}(R^4)\text{SO}_2\text{NH}_2$ ,  $-\text{N}(R^4)\text{SO}_2\text{NHR}^7$ ,  $-\text{N}(R^4)\text{SO}_2\text{N}(R^7)_2$ ,  $-\text{N}(R^4)\text{COR}^7$ ,  $-\text{N}(R^4)\text{CON}(R^7)_2$ ,  $-\text{N}(R^4)\text{CSN}(R^7)_2$ ,  $-\text{N}(R^4)\text{CSR}^7$ ,  $-\text{N}(R^4)\text{C(O)OR}^7$ ,  $-\text{SO}_2\text{NHet}^1$  [where  $-\text{NHet}^1$  is an optionally substituted  $\text{C}_{5-7}$ cyclicamino group optionally containing one or more other  $-\text{O}-$  or  $-\text{S}-$  atoms or  $-\text{N}(R^4)-$ ,  $-\text{C(O)}-$  or  $-\text{C(S)}-$  groups],  $-\text{CONHet}^1$ ,  $-\text{CSNHet}^1$ ,  $-\text{N}(R^4)\text{SO}_2\text{NHet}^1$ ,  $-\text{N}(R^4)\text{CONHet}^1$ ,  $-\text{N}(R^4)\text{CSNHet}^1$ ,  $-\text{SO}_2\text{N}(R^4)\text{Het}^2$  [where  $\text{Het}^2$  is an optionally substituted monocyclic  $\text{C}_{5-7}$ carbocyclic group optionally containing one or more  $-\text{O}-$  or  $-\text{S}-$  atoms or  $-\text{N}(R^4)-$ ,  $-\text{C(O)}-$  or  $-\text{C(S)}-$  groups],  $-\text{CON}(R^4)\text{Het}^2$ ,  $-\text{CSN}(R^4)\text{Het}^2$ ,  $-\text{N}(R^4)\text{CON}(R^4)\text{Het}^2$ ,  $-\text{N}(R^4)\text{CSN}(R^4)\text{Het}^2$ , aryl or heteroaryl group;  $\text{Alk}^3$  is a straight or branched  $\text{C}_{1-6}$ alkylene,  $\text{C}_{2-6}$ alkenylene or  $\text{C}_{2-6}$ alkynylene chain, optionally interrupted by one, two or three  $-\text{O}-$  or  $-\text{S}-$  atoms or  $-\text{S(O)}_n$  [where  $n$  is an integer 1 or 2] or  $-\text{N}(R^8)-$  groups [where  $R^8$  is a hydrogen atom or  $\text{C}_{1-6}$ alkyl, e.g. methyl or ethyl group]; and  $m$  is zero or an integer 1, 2 or 3. It will be appreciated that when two  $R^4$  or  $R^7$  groups are present in one of the above substituents, the  $R^4$  or  $R^7$  groups may be the same or different.
- When in the group  $-\text{Alk}^3(R^{6a})_m$   $m$  is an integer 1, 2 or 3, it is to be understood that the substituent or substituents  $R^{6a}$  may be present on any suitable carbon atom in  $-\text{Alk}^3$ . Where more than one  $R^{6a}$  substituent is present these may be the same or different and may be present on the same or different atom in  $-\text{Alk}^3$ . Clearly, when  $m$  is zero and no substituent  $R^{6a}$  is present the alkylene, alkenylene or alkynylene chain represented by  $\text{Alk}^3$  becomes an alkyl, alkenyl or alkynyl group.

When  $R^{6a}$  is a substituted amino group it may be for example a group  $-NHR^7$  [where  $R^7$  is as defined above] or a group  $-N(R^7)_2$  wherein each  $R^7$  group is the same or different.

5

When  $R^{6a}$  is a halogen atom it may be for example a fluorine, chlorine, bromine, or iodine atom.

10 When  $R^{6a}$  is a substituted hydroxyl or substituted thiol group it may be for example a group  $-OR^7$  or a  $-SR^7$  or  $-SC(=NH)NH_2$  group respectively.

15 Esterified carboxyl groups represented by the group  $R^{6a}$  include groups of formula  $-CO_2Alk^4$  wherein  $Alk^4$  is a straight or branched, optionally substituted  $C_{1-8}$ alkyl group such as a methyl, ethyl, n-propyl, i-propyl, n-butyl, i-butyl, s-butyl or t-butyl group; a  $C_{6-12}$ aryl $C_{1-8}$ alkyl group such as an optionally substituted benzyl, phenylethyl, phenylpropyl, 1-naphthylmethyl or 2-naphthylmethyl group; a  $C_{6-12}$ aryl group such as an optionally substituted phenyl, 1-naphthyl or 2-naphthyl group; a  $C_{6-12}$ aryloxy $C_{1-8}$ alkyl group such as an optionally substituted phenyloxymethyl, phenyloxyethyl, 1-naphthyl-oxymethyl, or 2-naphthyl-oxymethyl group; an optionally substituted  $C_{1-8}$ alkanoyloxy $C_{1-8}$ alkyl group, such as a pivaloyloxymethyl, propionyloxyethyl or propionyloxypropyl group; or a  $C_{6-12}$ aroyloxy $C_{1-8}$ alkyl group such as an optionally substituted benzoyloxyethyl or benzoyloxypropyl group. Optional substituents present on the  $Alk^4$  group include  $R^{6a}$  substituents described above.

20

25

When  $Alk^3$  is present in or as a substituent it may be for example a methylene, ethylene, n-propylene, i-propylene, n-butylene, i-butylene, s-butylene, t-butylene, ethenylene, 2-propenylene, 2-butenylene, 3-butenylene, ethynylene, 2-propynylene, 2-butynylene or 3-butynylene chain, optionally interrupted by one, two, or three  $-O-$  or  $-S-$ , atoms or  $-S(O)-$ ,  $-S(O)_2-$  or  $-N(R^8)-$  groups.

30

35 Aryl or heteroaryl groups represented by the groups  $R^{6a}$  or  $R^7$  include mono- or bicyclic optionally substituted  $C_{6-12}$  aromatic or  $C_{1-9}$  heteroaromatic groups as described above for the groups  $R^1$  and Het.

The aromatic and heteroaromatic groups may be attached to the remainder of the compound of formula (1) by any carbon or hetero e.g. nitrogen atom as appropriate.

- 5 When -NHet<sup>1</sup> or -Het<sup>2</sup> forms part of a substituent R<sup>6</sup> each may be for example an optionally substituted pyrrolidinyl, pyrazolidinyl, piperazinyl, morpholinyl, thiomorpholinyl, piperidinyl or thiazolidinyl group. Additionally Het<sup>2</sup> may represent for example, an optionally substituted cyclopentyl or cyclohexyl group. Optional substituents which may be present on -NHet<sup>1</sup>  
 10 or -Het<sup>2</sup> include those R<sup>5</sup> substituents described above.

- Particularly useful atoms or groups represented by R<sup>6</sup> include fluorine, chlorine, bromine or iodine atoms, or C<sub>1-6</sub>alkyl, e.g. methyl, ethyl, n-propyl, i-propyl, n-butyl or t-butyl, optionally substituted phenyl, pyridyl, pyrrolyl,  
 15 furyl, thiazolyl, or thienyl, C<sub>1-6</sub>alkylamino, e.g. methylamino or ethylamino, C<sub>1-6</sub>hydroxyalkyl, e.g. hydroxymethyl or hydroxyethyl, carboxyC<sub>1-6</sub>alkyl, e.g. carboxyethyl, C<sub>1-6</sub>alkylthio e.g. methylthio or ethylthio, carboxyC<sub>1-6</sub>alkylthio, e.g. carboxymethylthio, 2-carboxyethylthio or 3-carboxypropylthio, C<sub>1-6</sub>alkoxy, e.g. methoxy or ethoxy, hydroxyC<sub>1-6</sub>alkoxy, e.g. 2-  
 20 hydroxyethoxy, optionally substituted phenoxy, pyridyloxy, thiazolyloxy, phenylthio or pyridylthio, C<sub>5-7</sub>cycloalkoxy, e.g. cyclopentyloxy, haloC<sub>1-6</sub>alkyl, e.g. trifluoromethyl, haloC<sub>1-6</sub>alkoxy, e.g. trifluoromethoxy, C<sub>1-6</sub>alkylamino, e.g. methylamino or ethylamino, amino (-NH<sub>2</sub>), aminoC<sub>1-6</sub>alkyl, e.g. aminomethyl or aminoethyl, C<sub>1-6</sub>dialkylamino, e.g.  
 25 dimethylamino or diethylamino, C<sub>1-6</sub>alkylaminoC<sub>1-6</sub>alkyl, e.g. ethylaminoethyl, C<sub>1-6</sub>dialkylaminoC<sub>1-6</sub>alkyl, e.g. diethylaminoethyl, aminoC<sub>1-6</sub>alkoxy, e.g. aminoethoxy, C<sub>1-6</sub>alkylaminoC<sub>1-6</sub>alkoxy, e.g. methylaminoethoxy, C<sub>1-6</sub>dialkylaminoC<sub>1-6</sub>alkoxy, e.g. dimethylaminoethoxy, diethylaminoethoxy, isopropylaminoethoxy, or dimethylaminopropoxy, imido, such as  
 30 phthalimido or naphthalimido, e.g. 1,8-naphthalimido, nitro, cyano, amidino, hydroxyl (-OH), formyl [HC(O)-], carboxyl (-CO<sub>2</sub>H), -CO<sub>2</sub>Alk<sup>4</sup> [where Alk<sup>4</sup> is as defined above], C<sub>1-6</sub>alkanoyl e.g. acetyl, optionally substituted benzoyl, thiol (-SH), thioC<sub>1-6</sub>alkyl, e.g. thiomethyl or thioethyl, -SC(=NH)NH<sub>2</sub>, sulphonyl (-SO<sub>3</sub>H), C<sub>1-6</sub>alkylsulphonyl, e.g. methyl-  
 35 sulphonyl, optionally substituted phenylsulphonyl, aminosulphonyl (-SO<sub>2</sub>NH<sub>2</sub>), C<sub>1-6</sub>alkylaminosulphonyl, e.g. methylaminosulphonyl or

- ethylaminosulphonyl, C<sub>1-6</sub>dialkylaminosulphonyl, e.g. dimethylaminosulphonyl or diethylaminosulphonyl, phenylamino-sulphonyl, carboxamido (-CONH<sub>2</sub>), C<sub>1-6</sub>alkylaminocarbonyl, e.g. methylaminocarbonyl or ethylaminocarbonyl, C<sub>1-6</sub>dialkylaminocarbonyl, e.g. dimethylaminocarbonyl or diethylaminocarbonyl, aminoC<sub>1-6</sub>alkylaminocarbonyl, e.g. aminoethylaminocarbonyl, C<sub>1-6</sub>dialkylaminoC<sub>1-6</sub>alkylaminocarbonyl, e.g. diethylaminoethylaminocarbonyl, aminocarbonylamino, C<sub>1-6</sub>alkylaminocarbonylamino, e.g. methylaminocarbonylamino or ethylaminocarbonylamino, C<sub>1-6</sub>dialkylaminocarbonylamino, e.g. dimethylaminocarbonylamino or diethylaminocarbonylamino, C<sub>1-6</sub>alkylaminocarbonylC<sub>1-6</sub>alkylamino, e.g. methylaminocarbonylmethylamino, aminothiocabonylamino, C<sub>1-6</sub>alkylaminothiocabonylamino, e.g. methylaminothiocabonylamino or ethylaminothiocabonylamino, C<sub>1-6</sub>dialkylaminothiocabonylamino, e.g. dimethylaminothiocabonylamino or diethylaminothiocabonylamino, C<sub>1-6</sub>alkylaminothiocabonylC<sub>1-6</sub>alkylamino, e.g. ethylaminothiocabonylmethylamino, -CONHC(=NH)NH<sub>2</sub>, C<sub>1-6</sub>alkylsulphonylamino, e.g. methylsulphonylamino or ethylsulphonylamino, C<sub>1-6</sub>dialkylsulphonylamino, e.g. dimethylsulphonylamino or diethylsulphonylamino, optionally substituted phenylsulphonylamino, aminosulphonylamino (-NHSO<sub>2</sub>NH<sub>2</sub>), C<sub>1-6</sub>alkylaminosulphonylamino, e.g. methylaminosulphonylamino or ethylaminosulphonylamino, C<sub>1-6</sub>dialkylaminosulphonylamino, e.g. dimethylaminosulphonylamino or diethylaminosulphonylamino, optionally substituted morpholinesulphonylamino or morpholinesulphonylC<sub>1-6</sub>alkylamino, optionally substituted phenylaminosulphonylamino, C<sub>1-6</sub>alkanoylamino, e.g. acetylamino, aminoC<sub>1-6</sub>alkanoylamino e.g. aminoacetylamino, C<sub>1-6</sub>dialkylaminoC<sub>1-6</sub>alkanoylamino, e.g. dimethylaminoacetylamino, C<sub>1-6</sub>alkanoylaminoC<sub>1-6</sub>alkyl, e.g. acetylaminomethyl, C<sub>1-6</sub>alkanoylaminoC<sub>1-6</sub>alkylamino, e.g. acetamidoethylamino, C<sub>1-6</sub>alkoxycarbonylamino, e.g. methoxycarbonylamino, ethoxycarbonylamino or t-butoxycarbonylamino or optionally substituted benzyloxy, pyridylmethoxy, thiazolylmethoxy, benzyloxycarbonylamino, benzyloxycarbonylaminoC<sub>1-6</sub>alkyl e.g. benzyloxycarbonylaminoethyl, benzylthio, pyridylmethylthio or thiazolylmethylthio groups.
- In the above groups of particularly useful R<sup>6</sup> substituents, the reference to optional substitution is intended to relate primarily to the aromatic or

heteroaromatic portions of the groups described. Thus for example such groups may be optionally mono-, di- or tri-substituted by those particular atoms or groups described above for each of  $R^a$  and  $R^b$ .

- 5 Where desired, two  $R^6$  substituents may be linked together to form a cyclic group such as a cyclic ether, e.g. a  $C_{1-6}$ alkylenedioxy group such as methylenedioxy or ethylenedioxy.

- 10 It will be appreciated that where two or more  $R^6$  substituents are present, these need not necessarily be the same atoms and/or groups. In general, the substituent(s) may be present at any available ring position in the heteroaromatic group represented by Het.

- 15 The presence of certain substituents in the compounds of formula (1) may enable salts of the compounds to be formed. Suitable salts include pharmaceutically acceptable salts, for example acid addition salts derived from inorganic or organic acids, and salts derived from inorganic and organic bases.

- 20 Acid addition salts include hydrochlorides, hydrobromides, hydroiodides, alkylsulphonates, e.g. methanesulphonates, ethanesulphonates, or isothionates, arylsulphonates, e.g. p-toluenesulphonates, besylates or napsylates, phosphates, sulphates, hydrogen sulphates, acetates, trifluoroacetates, propionates, citrates, maleates, fumarates, malonates, 25 succinates, lactates, oxalates, tartrates and benzoates.

- Salts derived from inorganic or organic bases include alkali metal salts such as sodium or potassium salts, alkaline earth metal salts such as magnesium or calcium salts, and organic amine salts such as morpholine, 30 piperidine, dimethylamine or diethylamine salts.

- Particularly useful salts of compounds according to the invention include pharmaceutically acceptable salts, especially acid addition pharmaceutically acceptable salts.

- 35 R in compounds of the invention is preferably a  $-CO_2H$  group.

When present, the aliphatic chain represented by  $\text{Alk}^1$  in compounds of the invention is preferably a  $-\text{CH}_2-$  chain.

5  $\text{Alk}^2$  in compounds of formula (1) is preferably a  $-\text{CH}_2-$  chain and  $m$  is preferably an integer 1. In compounds of this type, the carbon atom to which  $\text{Alk}^2$  and  $\text{R}$  are attached forms a chiral centre and is preferably in the L configuration.

10  $\text{R}^2$  in compounds of formula (1) is preferably a hydrogen atom.

$\text{R}^3$  in compounds of the invention is preferably a hydrogen atom.

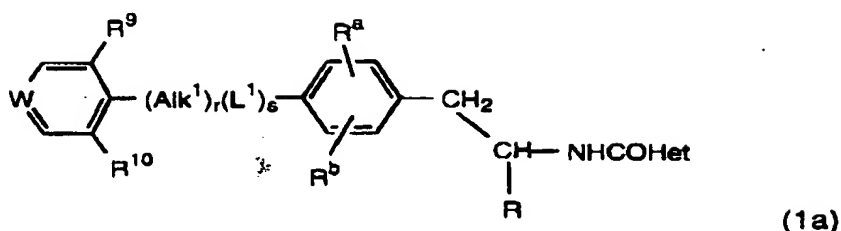
15 In general in compounds of the invention  $-(\text{Alk}^1)_r(\text{L}^1)_s$  is preferably  $-\text{CH}_2\text{O}-$ ,  $-\text{SO}_2\text{NH}-$ ,  $-\text{C}(\text{O})\text{O}-$  or  $-\text{CON}(\text{R}^4)-$  and is especially  $-\text{CONH}-$ .

20 In general in compounds of the invention the group  $\text{R}^1$  is preferably an optionally substituted aromatic or heteroaromatic group. Particularly useful groups of these types include optionally substituted phenyl, pyridyl or pyrimidinyl groups, particularly those in which the substituent when present is an atom or group  $-\text{L}^2(\text{CH}_2)_p\text{L}^3(\text{R}^c)_q$  as described above. Each substituent may be present on any available ring carbon or nitrogen atom.

25 The heteroaromatic group represented by  $\text{Het}$  in compounds of formula (1) is preferably an optionally substituted  $\text{C}_{3-5}$  monocyclic heteroaromatic group containing one, two or three heteroatoms selected from oxygen, sulphur or nitrogen atoms. Particularly useful groups of this type include optionally substituted pyrrolyl and pyridyl groups. Especially useful heteroaromatic groups represented by  $\text{Het}$  include optionally substituted 3- or 4-pyridyl groups, particularly 2-monosubstituted 3- or 4-pyridyl or 2,6-disubstituted 3- or 4-pyridyl groups. In these, and in general in the group  $\text{Het}$ , the optional substituent when present is preferably an atom or group  $\text{R}^6$  as defined above.

35 A particularly useful class of compounds according to the invention has the formula (1a)





5 wherein -W= is -CH= or -N=,  $R^9$  and  $R^{10}$ , which may be the same or different is each a  $-L^2(CH_2)_pL^3(R^c)_q$  atom or group as generally and particularly defined above, and  $Alk^1$ ,  $r$ ,  $L^1$ ,  $s$ ,  $R^a$ ,  $R^b$ ,  $R$  and Het are as generally and particularly defined above, and the salts, solvates, hydrates and N-oxides thereof.

10 It will be appreciated that the various preferences stated above in relation to groups present in compounds of formula (1) apply equally to the same groups when present in compounds of formula (1).

15 Additionally, in the compounds of formula (1a)  $-(Alk^1)_r(L^1)_s-$  is preferably a  $-CH_2O$  or  $-CON(R^4)-$  group and is especially a  $-CONH-$  group. Het is preferably an optionally substituted pyrrolyl or especially an optionally substituted pyridyl group.

20 Particularly useful compounds of formula (1a) are those wherein Het is a 2-monosubstituted 3- or 4-pyridyl group or a 2,6-disubstituted 3- or 4-pyridyl group.

25 One of  $R^9$  or  $R^{10}$  in compounds of formula (1a) may be for example a hydrogen atom and the other a substituent  $L^2(CH_2)_pL^3(R^c)_q$  in which  $R^c$  is not a covalent bond and  $p$  is zero, but preferably each of  $R^9$  and  $R^{10}$  is a substituent  $-L^2(CH_2)_pL^3(R^c)_q$  where  $R^c$  is as just defined. Particularly useful  $R^9$  or  $R^{10}$  substituents include a hydrogen atom or halogen atom, especially fluorine or chlorine atoms, or a methyl, ethyl, methoxy, ethoxy,  $-CF_3$ ,  $-OH$ ,  $-CN$ ,  $-NO_2$ ,  $-NH_2$ ,  $-NHCH_2$ ,  $-N(CH_3)_2$ ,  $-COCH_3$ ,  $-SCH_3$ ,  $-CO_2H$  or  $-CO_2CH_3$  group.

30

Particularly useful compounds according to the invention include the following:

2-Thio(S-2,5-dimethoxyphenyl)nicotinoyl-(*N*-2,6-dichlorobenzoyl)-*L*-4-aminophenylalanine;

2-Thio(S-2,5-dimethoxyphenyl)nicotinoyl-(*N*-2,6-dichlorobenzoyl)-*L*-4-aminophenylalanine;

5 *N*-(3,5-Dichloro-4-picolyl)-*N'*-(3,5-dichloro-4-picolyl)-*L*-4-amino-phenylalanine;

*N*-(2-Chloronicotinoyl)-*N'*-(3,5-dichloro-4-picolyl)-*L*-4-amino-phenylalanine;

*O*-(2,6-dichlorobenzyl)-*N*-(4-acetyl-1,2,5-trimethyl-3-pyrrolyl)-*L*-tyrosine;

10 (*N'*-3,5-Dichloroisonicotinoyl)-*N*-{[3-pyridinylmethyl]thio}isonicotinoyl)-*L*-4-aminophenylalanine;

*N*-(4-Acetyl-1,2,5-trimethyl-1*H*-pyrrole-3-carbonyl)-*N'*-(3,5-dichloro-4-picolyl)-*L*-4-aminophenylalanine; and the salts, solvates, hydrates and *N*-oxides thereof.

15 Compounds according to the invention are potent and selective inhibitors of  $\alpha_4$  integrins. The ability of the compounds to act in this way may be simply determined by employing tests such as those described in the Examples hereinafter. In particular compounds of the invention, such as  
20 the compounds of formula (1a) herein, the compounds are advantageously selective  $\alpha_4\beta_1$  inhibitors.

The compounds are of use in modulating cell adhesion and in particular are of use in the prophylaxis and treatment of diseases or disorders involving inflammation in which the extravasation of leukocytes plays a role  
25 and the invention extends to such a use and to the use of the compounds for the manufacture of a medicament for treating such diseases or disorders.

30 Diseases or disorders of this type include inflammatory arthritis such as rheumatoid arthritis vasculitis or polydermatomyositis, multiple sclerosis, allograft rejection, diabetes, inflammatory dermatoses such as psoriasis or dermatitis, asthma and inflammatory bowel disease.

35 For the prophylaxis or treatment of disease the compounds according to the invention may be administered as pharmaceutical compositions, and according to a further aspect of the invention we provide a pharmaceutical

composition which comprises a compound of formula (1) together with one or more pharmaceutically acceptable carriers, excipients or diluents.

- 5      Pharmaceutical compositions according to the invention may take a form suitable for oral, buccal, parenteral, nasal, topical or rectal administration, or a form suitable for administration by inhalation or insufflation.

- 10      For oral administration, the pharmaceutical compositions may take the form of, for example, tablets, lozenges or capsules prepared by conventional means with pharmaceutically acceptable excipients such as binding agents (e.g. pregelatinised maize starch, polyvinylpyrrolidone or hydroxypropyl methylcellulose); fillers (e.g. lactose, microcrystalline cellulose or calcium hydrogen phosphate); lubricants (e.g. magnesium stearate, talc or silica); disintegrants (e.g. potato starch or sodium glycollate); or wetting agents (e.g. sodium lauryl sulphate). The tablets  
15      may be coated by methods well known in the art. Liquid preparations for oral administration may take the form of, for example, solutions, syrups or suspensions, or they may be presented as a dry product for constitution with water or other suitable vehicle before use. Such liquid preparations  
20      may be prepared by conventional means with pharmaceutically acceptable additives such as suspending agents, emulsifying agents, non-aqueous vehicles and preservatives. The preparations may also contain buffer salts, flavouring, colouring and sweetening agents as appropriate.
- 25      Preparations for oral administration may be suitably formulated to give controlled release of the active compound.

- For buccal administration the compositions may take the form of tablets or lozenges formulated in conventional manner.

- 30      The compounds for formula (1) may be formulated for parenteral administration by injection e.g. by bolus injection or infusion. Formulations for injection may be presented in unit dosage form, e.g. in glass ampoule or multi dose containers, e.g. glass vials. The compositions for injection  
35      may take such forms as suspensions, solutions or emulsions in oily or aqueous vehicles, and may contain formulatory agents such as

suspending, stabilising, preserving and/or dispersing agents. Alternatively, the active ingredient may be in powder form for constitution with a suitable vehicle, e.g. sterile pyrogen-free water, before use.

5 In addition to the formulations described above, the compounds of formula (1) may also be formulated as a depot preparation. Such long acting formulations may be administered by implantation or by intramuscular injection.

10 For nasal administration or administration by inhalation, the compounds for use according to the present invention are conveniently delivered in the form of an aerosol spray presentation for pressurised packs or a nebuliser, with the use of suitable propellant, e.g. dichlorodifluoromethane, trichlorofluoromethane, dichlorotetrafluoroethane, carbon dioxide or other suitable  
15 gas or mixture of gases.

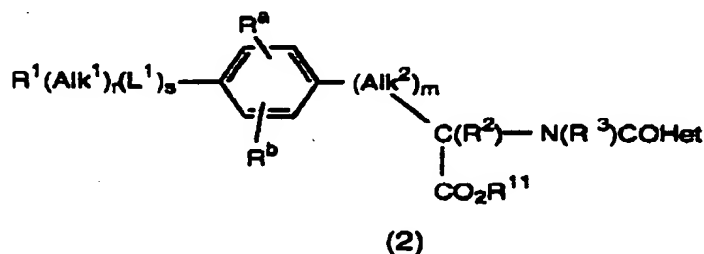
The compositions may, if desired, be presented in a pack or dispenser device which may contain one or more unit dosage forms containing the active ingredient. The pack or dispensing device may be accompanied by  
20 instructions for administration.

The quantity of a compound of the invention required for the prophylaxis or treatment of a particular condition will vary depending on the compound chosen, and the condition of the patient to be treated. In general,  
25 however, daily dosages may range from around 100ng/kg to 100mg/kg e.g. around 0.01mg/kg to 40mg/kg body weight for oral or buccal administration, from around 10ng/kg to 50mg/kg body weight for parenteral administration and around 0.05mg to around 1000mg e.g. around 0.5mg to around 1000mg for nasal administration or administration by inhalation  
30 or insufflation.

The compounds of the invention may be prepared by a number of processes as generally described below and more specifically in the Examples hereinafter. In the following process description, the symbols  
35 R, R<sup>1</sup>-R<sup>3</sup>, R<sup>a</sup>, R<sup>b</sup>, L<sup>1</sup>, Alk<sup>1</sup>, Alk<sup>2</sup>, m, r, s and Het when used in the formulae depicted are to be understood to represent those groups described above

in relation to formula (1) unless otherwise indicated. In the reactions described below, it may be necessary to protect reactive functional groups, for example hydroxy, amino, thio or carboxy groups, where these are desired in the final product, to avoid their unwanted participation in the reactions. Conventional protecting groups may be used in accordance with standard practice [see, for example, Green, T. W. in "Protective Groups in Organic Synthesis", John Wiley and Sons, 1991]. In some instances, deprotection may be the final step in the synthesis of a compound of formula (1) and the processes according to the invention described hereinafter are to be understood to extend to such removal of protecting groups.

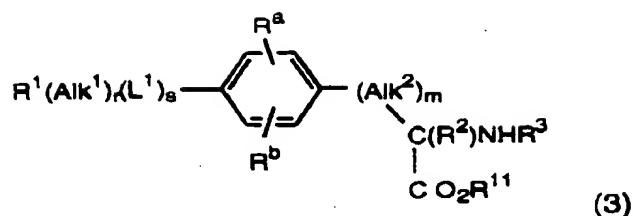
Thus according to a further aspect of the invention, a compound of formula (1) may be obtained by hydrolysis of an ester of formula (2):



where R<sup>11</sup> is an alkyl group.

The hydrolysis may be performed using either an acid or a base depending on the nature of R<sup>11</sup>, for example an organic acid such as trifluoroacetic acid or an inorganic base such as lithium hydroxide optionally in an aqueous organic solvent such as an amide, e.g. a substituted amide such as dimethylformamide, an ether, e.g. a cyclic ether such as tetrahydrofuran or dioxane or an alcohol, e.g. methanol at around ambient temperature. Where desired, mixtures of such solvents may be used.

Esters of formula (2) may be prepared by coupling an amine of formula (3):



(where  $\text{R}^{11}$  is as just described) or a salt thereof with an acid of formula (4):

5



or an active derivative thereof.

- 10 Active derivatives of acids of formula (4) include anhydrides, esters and halides. Particular esters include pentafluorophenyl or succinyl esters.

15 The coupling reaction may be performed using standard conditions for reactions of this type. Thus for example the reaction may be carried out in a solvent, for example an inert organic solvent such as an amide, e.g. a substituted amide such as dimethylformamide, an ether, e.g. a cyclic ether such as tetrahydrofuran, or a halogenated hydrocarbon, such as dichloromethane, at a low temperature, e.g. around  $-30^\circ\text{C}$  to around ambient temperature, optionally in the presence of a base, e.g. an organic

20 base such as an amine, e.g. triethylamine, pyridine, or dimethylaminopyridine, or a cyclic amine, such as N-methylmorpholine.

25 Where an acid of formula (4) is used, the reaction may additionally be performed in the presence of a condensing agent, for example a diimide such as 1-(3-dimethylaminopropyl)-3-ethylcarbodiimide or N,N'-dicyclohexylcarbodiimide, advantageously in the presence of a catalyst such as a N-hydroxy compound e.g. a N-hydroxytriazole such as 1-hydroxybenzotriazole. Alternatively, the acid may be reacted with a chloroformate, for example ethylchloroformate, prior to reaction with the amine of formula

30 (3).

Intermediates of formulae (2), (3) and (4), or compounds of formula (1), may be manipulated to introduce substituents to aromatic or heteroaromatic groups or modify existing substituents in groups of these types. Typically, such manipulation may involve standard substitution approaches employing for example alkylation, arylation, heteroarylation, acylation, thioacylation, halogenation, sulphonylation, nitration, formylation or coupling reactions. Alternatively, existing substituents may be modified for example by oxidation, reduction or cleavage reactions. Particular examples of such reactions are given below. Where these are described in relation to the generation of the group  $R^1(Alk^1)_r(L^1)_s-$ , it will be appreciated that each reaction may also be used to introduce or modify  $R^5$  and/or  $R^6$  substituents as appropriate.

Thus in one example, a compound wherein  $R^1(Alk^1)_r(L^1)_s-$  is a  $-L^1H$  group may be alkylated, arylated or heteroarylated using a reagent  $R^1(Alk^1)_rX$  in which  $R^1$  is other than a hydrogen atom and  $X$  is a leaving atom or group such as a halogen atom, e.g. a fluorine, bromine, iodine or chlorine atom or a sulphonyloxy group such as an alkylsulphonyloxy, e.g. trifluoromethylsulphonyloxy or arylsulphonyloxy, e.g. p-toluenesulphonyloxy group.

The reaction may be carried out in the presence of a base such as a carbonate, e.g. caesium or potassium carbonate, an alkoxide, e.g. potassium t-butoxide, or a hydride, e.g. sodium hydride, in a dipolar aprotic solvent such as an amide, e.g. a substituted amide such as dimethylformamide or an ether, e.g. a cyclic ether such as tetrahydrofuran.

In another example, a compound where  $R^1(Alk^1)_r(L^1)_s$  is a  $-L^1H$  group is a hydrogen atom may be functionalised by acylation or thioacylation, for example by reaction with a reagent  $R^1(Alk^1)_rL^1X$  [wherein  $L^1$  is a  $-C(O)-$ ,  $C(S)-$ ,  $-N(R^4)C(O)-$  or  $N(R^4)C(S)-$  group], in the presence of a base, such as a hydride, e.g. sodium hydride or an amine, e.g. triethylamine or N-methylmorpholine, in a solvent such as a halogenated hydrocarbon, e.g. dichloromethane or carbon tetrachloride or an amide, e.g. dimethylformamide, at for example ambient temperature, or by reaction

with  $R^1(Alk^1)_rCO_2H$ ,  $R^1(Alk^1)_4COSH$  or an activated derivative thereof, for example as described above for the preparation of esters of formula (2).

5 In a further example a compound may be obtained by sulphonylation of a compound where  $R^1(Alk^1)_r(L^1)_s$  is an  $-OH$  group by reaction with a reagent  $R^1(Alk^1)_rL^1Hal$  [in which  $L^1$  is  $-S(O)-$  or  $-SO_2-$  and  $Hal$  is a halogen atom such as chlorine atom] in the presence of a base, for example an inorganic base such as sodium hydride in a solvent such as an amide, e.g. a substituted amide such as dimethylformamide at for example ambient  
10 temperature.

In another example, a compound where  $R^1(Alk^1)_r(L^1)_s$  is a  $-L^1H$  group, may be coupled with a reagent  $R^1OH$  (where  $R^1$  is other than a hydrogen atom) or  $R^1Alk^1OH$  in a solvent such as tetrahydrofuran in the presence of  
15 a phosphine, e.g. triphenylphosphine and an activator such as diethyl, diisopropyl- or dimethylazodicarboxylate to yield a compound containing a  $R^1(Alk^1)_rO-$  group.

In a further example, ester groups  $-CO_2R^4$  or  $-CO_2Alk^4$  in compounds of  
20 formula (1) may be converted to the corresponding acid  $[-CO_2H]$  by acid- or base-catalysed hydrolysis depending on the nature of the group  $R^4$  or  $Alk^4$ . Acid- or base-catalysed hydrolysis may be achieved for example by treatment with an organic or inorganic acid, e.g. trifluoroacetic acid in an aqueous solvent or a mineral acid such as hydrochloric acid in a solvent  
25 such as dioxan or an alkali metal hydroxide, e.g. lithium hydroxide in an aqueous alcohol, e.g. aqueous methanol.

In a second example,  $-OR^7$  [where  $R^7$  represents an alkyl group such as methyl group] groups in compounds of formula (1) may be cleaved to the  
30 corresponding alcohol  $-OH$  by reaction with boron tribromide in a solvent such as a halogenated hydrocarbon, e.g. dichloromethane at a low temperature, e.g. around  $-78^\circ C$ .

Alcohol  $[-OH]$  groups may also be obtained by hydrogenation of a  
35 corresponding  $-OCH_2R^7$  group (where  $R^7$  is an aryl group) using a metal catalyst, for example palladium on a support such as carbon in a solvent



such as ethanol in the presence of ammonium formate, cyclohexadiene or hydrogen, from around ambient to the reflux temperature. In another example, -OH groups may be generated from the corresponding ester [-CO<sub>2</sub>Alk<sup>4</sup> or CO<sub>2</sub>R<sup>4</sup>] or aldehyde [-CHO] by reduction, using for example a  
5 complex metal hydride such as lithium aluminium hydride or sodium borohydride in a solvent such as methanol.

In another example, alcohol -OH groups in compounds of formula (1) may be converted to a corresponding -OR<sup>3</sup> group by coupling with a reagent  
10 R<sup>7</sup>OH in a solvent such as tetrahydrofuran in the presence of a phosphine, e.g. triphenylphosphine and an activator such as diethyl-, diisopropyl-, or dimethylazodicarboxylate.

Aminosulphonylamino [-NHSO<sub>2</sub>NH<sub>2</sub>] groups in compounds of formula (1)  
15 may be obtained, in another example, by reaction of a corresponding amine [-NH<sub>2</sub>] with sulphamide in the presence of an organic base such as pyridine at an elevated temperature, e.g. the reflux temperature.

In a further example amine (-NH<sub>2</sub>) groups may be alkylated using a  
20 reductive alkylation process employing an aldehyde and a borohydride, for example sodium triacetoxyborohydride or sodium cyanoborohydride, in a solvent such as a halogenated hydrocarbon, e.g. dichloromethane, a ketone such as acetone, or an alcohol, e.g. ethanol, where necessary in the presence of an acid such as acetic acid at around ambient  
25 temperature.

In a further example, amine [-NH<sub>2</sub>] groups in compounds of formula (1) may be obtained by hydrolysis from a corresponding imide by reaction with  
30 hydrazine in a solvent such as an alcohol, e.g. ethanol at ambient temperature.

In another example, a nitro [-NO<sub>2</sub>] group may be reduced to an amine [-NH<sub>2</sub>], for example by catalytic hydrogenation using for example hydrogen  
35 in the presence of a metal catalyst, for example palladium on a support such as carbon in a solvent such as an ether, e.g. tetrahydrofuran or an

alcohol e.g. methanol, or by chemical reduction using for example a metal, e.g. tin or iron, in the presence of an acid such as hydrochloric acid.

5 Aromatic halogen substituents in compounds of the invention may be subjected to halogen-metal exchange with a base, for example a lithium base such as n-butyl or t-butyl lithium, optionally at a low temperature, e.g. around  $-78^{\circ}\text{C}$ , in a solvent such as tetrahydrofuran and then quenched with an electrophile to introduce a desired substituent. Thus, for example, a formyl group may be introduced by using dimethylformamide as the  
10 electrophile; a thiomethyl group may be introduced by using dimethyldisulphide as the electrophile.

In another example, sulphur atoms in compounds of the invention, for example when present in the linker group  $\text{L}^1$  may be oxidised to the  
15 corresponding sulfoxide using an oxidising agent such as a peroxy acid, e.g. 3-chloroperoxybenzoic acid, in an inert solvent such as a halogenated hydrocarbon, e.g. dichloromethane, at around ambient temperature.

Intermediates of formulae (3) and (4),  $\text{R}^1(\text{Alk}^1)_r\text{X}$ ,  $\text{R}^1(\text{Alk}^1)_r\text{L}^1\text{X}$ ,  
20  $\text{R}^1(\text{Alk}^1)_r\text{CO}_2\text{H}$ ,  $\text{R}^1\text{OH}$  and  $\text{R}^1\text{Alk}^1\text{OH}$  are either known compounds or may be prepared from known starting materials by use of analogous processes to those used for the preparation of the known compounds and/or by treating known compounds by one or more of the alkylation, acylation and other manipulations described herein, such as particularly  
25 described for the preparation of the Intermediates in the exemplification selection hereinafter.

N-oxides of compounds of formula (1) may be prepared for example by oxidation of the corresponding nitrogen base using an oxidising agent such  
30 as hydrogen peroxide in the presence of an acid such as acetic acid, at an elevated temperature, for example around  $70^{\circ}\text{C}$  to  $80^{\circ}\text{C}$ , or alternatively by reaction with a peracid such as peracetic acid in a solvent, e.g. dichloromethane, at ambient temperature.

35 Salts of compounds of formula (1) may be prepared by reaction of a compound of formula (1) with an appropriate base in a suitable solvent or

mixture of solvents e.g. an organic solvent such as an ether e.g. diethylether, or an alcohol, e.g. ethanol using conventional procedures.

Where it is desired to obtain a particular enantiomer of a compound of formula (1) this may be produced from a corresponding mixture of enantiomers using any suitable conventional procedure for resolving enantiomers.

Thus for example diastereomeric derivatives, e.g. salts, may be produced by reaction of a mixture of enantiomers of formula (1) e.g. a racemate, and an appropriate chiral compound, e.g. a chiral base. The diastereomers may then be separated by any convenient means, for example by crystallisation and the desired enantiomer recovered, e.g. by treatment with an acid in the instance where the diastereomer is a salt.

In another resolution process a racemate of formula (1) may be separated using chiral High Performance Liquid Chromatography. Alternatively, if desired a particular enantiomer may be obtained by using an appropriate chiral intermediate in one of the processes described above.

The following Examples illustrate the invention. All temperatures are in °C. The following abbreviations are used:

EDC - 1-(3-dimethylaminopropyl)3-ethylcarbodiimide;	
DMF - dimethylformamide;	DMSO - dimethylsulphoxide;
HOBT - 1-hydroxybenzotriazole;	THF - tetrahydrofuran;
TFA - trifluoroacetic acid;	NMM - N-methylmorpholine;
DCM - dichloromethane;	Ph - phenyl;
BOC - <u>tert</u> -butoxycarbonyl;	EtOAc - ethyl acetate;
MeOH - methanol;	LDA - lithium diisopropylamide;
tyr - tyrosine;	Ar - aryl;
HetAr - heteroaryl;	pyr - pyridine;
thioprop - thiopropine;	Bu - butyl;
Me - methyl;	app - apparent

**INTERMEDIATE 1****2-Chloronicotinoyl-O-(2,6-dichlorobenzyl)-L-tyrosine methyl ester**

A solution of O-(2,6-dichlorobenzyl)-L-tyrosine methyl ester hydrochloride, (1.11g, 2.84mmol), 2-chloronicotinic acid (0.45g, 2.84mmol), EDC (0.60g, 3.13mmol), HOBT (0.46g, 3.41mmol) and NMM (0.467ml, 0.43g, 4.26mmol) in DCM (25ml) was stirred at room temperature for 24 h. The reaction mixture was partitioned between DCM (50ml) and 10% NaHCO<sub>3</sub> solution (30ml). The organic layer was separated, dried over MgSO<sub>4</sub> and the solvent removed under vacuum to give a pale yellow solid that was recrystallised from EtOAc/hexane to give the title compound as an off white solid (1.14g, 81%).  $\delta$ H (CDCl<sub>3</sub>) 8.46 (1H, dd,  $\downarrow$  2.0, 4.7Hz), 8.05 (1H, dd,  $\downarrow$  2.0, 7.6Hz), 7.34 (2H, m), 7.25 (2H, m), 7.11 (2H, m), 6.96 (3H, m), 5.24 (2H, s), 5.06 (1H, m), 3.80 (3H, s) and 3.24 (2H, m).

**15 INTERMEDIATE 2****2-Thio(S-2,5-dimethoxyphenyl)nicotinoyl-O-(2,6-dichlorobenzyl)-L-tyrosine methyl ester**

A solution of O-(2,6-dichlorobenzyl)-L-tyrosine methyl ester hydrochloride (0.78g, 2.0mmol), EDC (0.42g, 2.2mmol) HOBT (0.32g, 2.4mmol) and NMM (0.55ml, 0.50g, 5.0mmol) in DMF (10ml) was treated with a solution of 2-thio(2,5-dimethoxyphenyl)nicotinic acid (0.58g, 2.0mmol) in DMF (2ml) and stirred for 16h at room temperature. Solvent was removed *in vacuo* and the residue was partitioned between EtOAc (50ml) and 10% hydrochloric acid (25ml). The organic layer was separated, washed with 10% NaHCO<sub>3</sub> solution (30ml), dried over MgSO<sub>4</sub> and the solvent removed *in vacuo* to give a yellow oil which was purified by chromatography (SiO<sub>2</sub>; EtOAc/hexane 1:1) to give the title compound as a white foam (1.14g, 86%).  $\delta$ H (CDCl<sub>3</sub>) 8.39 (1H, dd,  $\downarrow$  1.9, 4.7Hz), 7.85 (1H, dd,  $\downarrow$  1.9, 7.7Hz), 7.47 (1H, d,  $\downarrow$  7.6Hz), 7.37-6.81 (11H, m), 5.18 (2H, s), 5.09 (1H, m), 3.78 (6H, s), 3.53 (3H, s) and 3.25 (2H, m).

**INTERMEDIATE 3****2-Mercaptonicotinoyl-O-(2,6-dichlorobenzyl)-L-tyrosine methyl ester**

A solution of O-(2,6-dichlorobenzyl)-L-tyrosine methyl ester hydrochloride (2.50g, 6.4mmol), 2-mercaptanicotinic acid (0.99g, 6.4mmol) and NMM (1.41ml, 1.29g, 12.8mmol) in DMF (10ml) was stirred at room temperature

for 64h. Solvent was removed *in vacuo* and the residue partitioned between DCM (30ml) and water (25ml). The aqueous layer was extracted with DCM (30ml) and the combined organic layers were washed with 10% NaHCO<sub>3</sub> solution (30ml), dried over MgSO<sub>4</sub> and the solvent removed *in vacuo* to give a brown oil which was purified by chromatography (SiO<sub>2</sub>; gradient elution, 4:1 EtOAc/hexane to 100% EtOAc) to give the title compound as a yellow foam, (2.92g, 93%).  $\delta$ H (CDCl<sub>3</sub>) 8.71 (1H, dd,  $\perp$  1.8, 7.6Hz), 8.05 (1H, s), 7.61 (1H, dd,  $\perp$  1.8, 6.1Hz), 7.35 (2H, m), 7.33-7.19 (2H, m), 6.94 (2H, d,  $\perp$  8.7Hz), 5.22 (2H, s), 4.97 (1H, m), 3.74 (3H, s) and 3.21 (2H, m).

#### **INTERMEDIATE 4**

##### **2-Thio(S-4-picolinylnicotinoyl-O-(2,6-dichlorobenzyl)-L-tyrosine methyl ester**

A solution of Intermediate 3 (0.50g, 1.0mmol) and 4-picolyl chloride hydrochloride (0.17g, 1.0mmol) in DCM (10ml) was treated with 1,8-diazabicyclo[5.4.0]undec-7-ene (0.31ml, 0.31g, 2.0mmol) and stirred at room temperature for 5h. The reaction was partitioned between water and DCM, the organic layer separated, dried over MgSO<sub>4</sub> and the solvent removed *in vacuo* to give a yellow gum that was purified by chromatography (SiO<sub>2</sub>, EtOAc), to give a pale yellow solid, which was recrystallised from EtOAc/hexane (1:1) to give the title compound as an off white solid (0.30g, 52%).  $\delta$ H (CDCl<sub>3</sub>) 8.47(2H, m), 7.69 (1H, dd,  $\perp$  1.8, 7.7Hz), 7.38-7.22 (6H, m), 7.06 (3H, m), 6.90 (2H, m), 6.62 (1H, d,  $\perp$  7.5Hz), 5.23 (2H, s), 5.03 (1H, m), 4.40 (2H, m), 3.79 (3H, s), 3.28 (1H, dd,  $\perp$  5.8, 14.1Hz) and 3.19 (1H, dd,  $\perp$  5.4, 14.1Hz).

#### **INTERMEDIATE 5**

##### **2-Thio(S-2,5-dimethoxyphenyl)nicotinoyl-L-4-aminophenylalanine methyl ester**

A solution of 4-amino-L-phenylalanine methyl ester dihydrochloride (0.53g, 2.0mmol), EDC (0.42g, 2.2mmol) HOBT (0.32g, 2.4mmol) and NMM (0.66ml, 0.61g, 6.0mmol) in DMF (10ml) was treated with a solution of 2-thio(2,5-dimethoxyphenyl)nicotinic acid (0.58g, 2.0mmol) in DMF (2ml) and stirred for 64h at room temperature. The solvent was removed *in vacuo*, and the residue partitioned between DCM (30ml) and water (20ml). The

organic layer was separated, washed with 10% NaHCO<sub>3</sub> (20ml) solution, dried over MgSO<sub>4</sub> and the solvent evaporated *in vacuo* to give a brown gum which was purified by chromatography (SiO<sub>2</sub>; EtOAc) to give the title compound as a yellow foam (0.67g, 72%).  $\delta$ H (CDCl<sub>3</sub>) 8.35 (1H, dd,  $\downarrow$  1.8, 4.8Hz), 7.81 (1H, dd,  $\downarrow$  1.9, 7.7Hz), 7.41 (1H, d,  $\downarrow$  7.6Hz), 7.12 (1H, d,  $\downarrow$  3.0Hz), 7.07-6.81 (5H, m), 6.49 (2H, d,  $\downarrow$  8.4Hz), 5.00 (1H, m), 3.76 (3H, s), 3.74 (3H, s), 3.55 (3H, s) and 3.14 (2H, m).

#### INTERMEDIATE 6

##### 10 2-Thio(S-2,5-dimethoxyphenyl)nicotinoyl-(N-2,6-dichlorobenzoyl)-L-4-aminophenylalanine methyl ester

A solution of Intermediate 5 (0.68g, 1.5mmol) and NMM (0.53ml, 0.49g, 4.8mmol) in DCM (20ml) was treated with 2,6-dichlorobenzoyl chloride (0.23ml, 0.33g, 1.6mmol) and the reaction stirred for 16h at room temperature, then partitioned between DCM (50ml) and 10% NaHCO<sub>3</sub> solution (30ml). The organic layer was separated, dried over MgSO<sub>4</sub> and the solvent evaporated *in vacuo* to give an off-white solid that was triturated with EtOAc/diethyl ether (2:1) to give the title compound as an off-white solid (0.44g, 46%).  $\delta$ H (MeOH-d<sub>4</sub>) 8.28 (1H, dd,  $\downarrow$  1.8, 4.9Hz), 7.70 (1H, dd,  $\downarrow$  1.8, 7.6Hz), 7.58 (2H, d,  $\downarrow$  8.6Hz), 7.48-7.38 (3H, m), 7.30 (2H, d,  $\downarrow$  8.6Hz), 7.27 (1H, dd,  $\downarrow$  4.9, 7.7Hz), 7.01 (1H, dd,  $\downarrow$  1.1, 2.3Hz), 6.93 (2H, m), 4.87 (1H, m), 3.77 (3H, s), 3.75 (3H, s), 3.60 (3H, s), 3.32 (1H, m) and 3.13 (1H, dd,  $\downarrow$  8.6, 14.0Hz).

#### 25 INTERMEDIATE 7

##### O-(2,6-dichlorobenzyl)-N-(4-acetyl-1,2,5-trimethyl-3-pyrrolyl)-L-tyrosine methyl ester

NMM (155mg, 169 $\mu$ l, 1.54mmol), HOBt (227mg, 1.68mmol), 4-acetyl-1,2,5-trimethylpyrrole-3-carboxylic acid (300mg, 1.54mmol) and EDC (295mg, 1.54mmol) were added sequentially to a stirred solution of O-(2,6-dichlorobenzyl)-L-tyrosine methyl ester hydrochloride (546mg, 1.40mmol) in dry DMF (15ml). The reaction was stirred at room temperature under N<sub>2</sub> for 18h. The solvent was removed *in vacuo* and the residue partitioned between EtOAc (50ml), and 10% aqueous Na<sub>2</sub>CO<sub>3</sub> (40ml). The phases were separated and the aqueous phase extracted with EtOAc (2 x 25ml). The combined organic extracts were washed consecutively with 5%

- aqueous hydrochloric acid (20ml), 10% aqueous Na<sub>2</sub>CO<sub>3</sub> (20ml) and brine (10ml), dried (Na<sub>2</sub>SO<sub>4</sub>) and evaporated *in vacuo*. The obtained orange foam (0.6g) was chromatographed (silica; 50% EtOAc/Hexane → 100% EtOAc) affording the title compound as a white foam (380mg, 51%); <sup>1</sup>Hnmr (d<sup>6</sup>DMSO) 8.53 (1H, d,  $\downarrow$  8Hz, NH), 7.57-7.43 (3H, m's, aryl-H), 7.22 (2H, d,  $\downarrow$  8.5Hz), aryl-H), 6.95 (2H, d,  $\downarrow$  8.5Hz, aryl-H), 5.18 (2H, br s, CH<sub>2</sub>-O), 4.67 (1H, m,  $\alpha$ -tyr-H), 3.66 (3H, s, Me-O), 3.34 (3H, s, Me-N), 3.12 (1H, dd,  $\downarrow$  4.1, 13.8Hz, CH<sub>A</sub>H<sub>B</sub>Ar), 2.90 (1H, dd,  $\downarrow$  11.3, 13.8Hz, CH<sub>A</sub>H<sub>B</sub>Ar), 2.32 (3H, s, MeCO), 2.0 (3H, s, pyrrole-Me) and 1.94 (3H, s, pyrrole-Me).  
 5  
 10 m/z (ES + 60V) 531 (MH<sup>+</sup>, 100), 533 (MH<sup>+</sup>, 75) 553 (MNa<sup>+</sup>, 15%).

### **INTERMEDIATE 8**

#### **2-Chloronicotiny-(N-2,6-dichlorobenzoyl)-L-4-aminophenylalanine methyl ester**

- 15 EDC (270mg, 1.5mmol) was added to a stirred solution of (N-2,6-dichlorobenzoyl)-L-4-aminophenylalanine methyl ester (500mg, 1.3mmol), 2 chloronicotinic acid (200mg, 1.3mmol), HOBT (190mg, 1.5mmol) and NMM (423 $\mu$ l, 3.9mmol) in anhydrous DMF (2ml) at 0°. The DMF solution was stirred overnight at room temperature then the DMF was evaporated  
 20 *in vacuo*. The residue was taken up in DCM (50ml), washed with water (3 x 10ml), saturated aqueous NaHCO<sub>3</sub> (2 x 10ml) and water (2 x 10ml), dried (Na<sub>2</sub>SO<sub>4</sub>) and evaporated *in vacuo*. The residue was purified by chromatography (SiO<sub>2</sub>; 1:1 EtOAc: hexane) to give the title compound as a white foam (450mg, 80%).  $\delta$ H (CDCl<sub>3</sub>), 8.50 (1H, m, pyr H), 8.05 (1H, d,  $\downarrow$  9.2Hz, pyrH), 7.58 (2H, d,  $\downarrow$  8.5Hz, ArH), 7.46 (1H, br s, NH), 7.36-7.24 (3H, m, 2ArH, 1pyrH), 7.21 (2H, d,  $\downarrow$  8.5Hz, 2 ArH), 7.12 (1H, d,  $\downarrow$  6.6Hz, NH), 5.05-5.18 (1H, m, CH $\alpha$ tyr), 3.81 (3H, s, CO<sub>2</sub>Me) and 3.39-3.18 (2H, m, CH<sub>2</sub>Ar).  
 25  
 30

### **INTERMEDIATE 9**

#### **Methyl-2-thio(S-acetate)nicotinoyl-O-(2,6-dichlorobenzyl)-L-tyrosine methyl ester**

- A solution of Intermediate 3 (370mg, 0.76mmol) in anhydrous DMF (2ml) was added to a suspension of sodium hydride (60% in oil, 33mg, 0.83mmol) in anhydrous DMF (3ml) at 0°. The mixture was stirred for  
 35 10min at room temperature, then recooled to 0°. Methyl bromoacetate

(115mg, 0.76mmol) was added dropwise, then the mixture was stirred overnight at room temperature. The mixture was quenched with water (0.5ml) and the DMF evaporated *in vacuo*. The residue was dissolved in EtOAc, washed with water (3 x 10ml), dried (Na<sub>2</sub>SO<sub>4</sub>) and evaporated *in vacuo*. The residue was purified by chromatography (SiO<sub>2</sub>; EtOAc/hexane 1:1) to give the title compound as a white solid (360mg, 85%).  $\delta$ H (CDCl<sub>3</sub>) 8.45 (1H, m, pyrH), 7.71 (1H, d,  $\downarrow$  7.7, pyrH), 7.35 (2H, d,  $\downarrow$  7.3Hz, 2ArH), 7.28-6.9 (6H, m, 5ArH, 1pyrH), 6.71 (1H, d,  $\downarrow$  7.5Hz, NH), 5.24 (2H, s, OCH<sub>2</sub>Ar), 5.08-5.03 (1H, m, CH $\alpha$  tyr), 3.96 (2H, s, SCH<sub>2</sub>), 3.78 (3H, s, CO<sub>2</sub>CH<sub>3</sub>), 3.71 (3H, s, CO<sub>2</sub>CH<sub>3</sub>), 3.31 (1H, dd,  $\downarrow$  14, 5.4Hz, CH<sub>A</sub>H<sub>B</sub>Ar) and 3.21 (1H, dd,  $\downarrow$  14, 5.2Hz, CH<sub>A</sub>H<sub>B</sub>Ar).

#### **INTERMEDIATE 10**

##### **2-Thio(S-methyl)nicotinoyl-O-(2,6-dichlorobenzyl)-L-tyrosine methyl ester**

EDC (540mg, 3mmol) was added to a stirred solution of (O-2,6-dichlorobenzyl)-L-tyrosine methyl ester (1g, 2.6mmol), 2 methylmercaptonicotinic acid (433mg, 2.6mmol), HOBT (364mg, 2.6mmol) and NMM (846 $\mu$ l, 7.8mmol) in anhydrous DMF (4ml) at 0°. The DMF solution was stirred overnight at room temperature, then the DMF was evaporated *in vacuo*. The residue was taken up in DCM (70ml), washed with water (3 x 15ml), saturated aqueous NaHCO<sub>3</sub> (2 x 15ml) and water (2 x 15ml), dried (Na<sub>2</sub>SO<sub>4</sub>) and evaporated *in vacuo*. The residue was purified by chromatography (SiO<sub>2</sub>; EtOAc) to give the title compound as a white solid (1.2gm, 92%).  $\delta$ H (CDCl<sub>3</sub>) 8.42 (1H, dd,  $\downarrow$  1.7, 4.8, 1pyrH), 7.67 (1H, dd,  $\downarrow$  1.7, 7.6, 1pyrH), 7.31 (2H, d,  $\downarrow$  8.5, ArH), 7.18 (1H, dd,  $\downarrow$  2.3, 8.5, ArH), 7.09 (3H, d, plus broad peak,  $\downarrow$  8.6, 2ArH, 1NH), 6.97 (1H, m, 1pyrH), 6.88 (2H, d,  $\downarrow$  8.6, 2ArH), 5.18 (2H, s, OCH<sub>2</sub>Ar), 4.96 (1H, m, CH $\alpha$ tyr), 3.72 (3H, s, OCH<sub>3</sub>), 3.25-3.04 (2H, m, CH<sub>2</sub>Ar) and 2.47 (3H, s, SCH<sub>3</sub>). m/z (ESI, GOU) 505 (MH<sup>+</sup>).

#### **INTERMEDIATE 11**

##### **Ethyl 3-(4-(((4-methoxybenzyl)oxy)carbonyl)phenyl)-2-((diphenylmethylene)amino)propanoate**

N-(Diphenylmethylene) glycine ethyl ester (6.6g, 24.6mmol) and potassium carbonate (6.8g, 49mmol) were added to a solution of 4-methoxybenzyl-4-



(bromomethyl)benzoate (8.2g, 24.6mmol) in acetonitrile (200ml). The mixture was heated at reflux overnight, then filtered and the solvent removed *in vacuo* to give the title compound as a yellow oil (13.55g).  $\delta$ H (CDCl<sub>3</sub>, 300 MHz) 7.8 (2H, d,  $\downarrow$  9.0Hz), 7.5 (10H, m), 7.3 (2H, d,  $\downarrow$  9.0Hz), 6.9 (2H, d), 6.6 (2H, m), 5.23 (2H, s), 4.1 (3H, m), 3.7 (3H, s), 3.2 (2H, m) and 1.2 (3H, m);  $m/z$  (ESI) 522 (MH<sup>+</sup>).

#### **INTERMEDIATE 12**

##### **Ethyl 2-amino-3-(4-((4-methoxybenzyl)oxy)carbonyl)phenyl)**

##### **propanoate**

Hydrochloric acid (2M, 15.83ml, 1.5eq) was added to a solution of Intermediate 11 (11.0g, 21.12mmol) in THF (30ml). After 20min the reaction mixture was basified to pH7 with NaHCO<sub>3</sub> and the solvent removed *in vacuo*. The residue was taken up in EtOAc (300ml) and washed with water (200ml) and brine (200ml), dried (MgSO<sub>4</sub>) and evaporated *in vacuo*. Chromatography (SiO<sub>2</sub>; EtOAc) gave the title compound as a yellow oil (4.91g, 65%).  $\delta$ H (CDCl<sub>3</sub>, 300MHz) 7.88 (2H, d,  $\downarrow$  8.0Hz), 7.4 (4H, dd), 7.0 (2H, d,  $\downarrow$  8.0Hz), 5.25 (2H, s), 4.05 (2H, q), 3.7 (3H, s), 3.57 (1H, t), 2.87 (2H, m) and 1.1 (3H, t),  $m/z$  (ESI) 358 (MH<sup>+</sup>).

#### **INTERMEDIATE 13**

##### **Ethyl 2-(((2-chloro-3-pyridinyl)carbonyl)amino)-3-(4-((4-methoxybenzyl)oxy)carbonyl)phenyl)propanoate**

EDC;HCl (591mg, 3.08mmol) and HOBt (416mg, 3.08mmol) were added to a solution of Intermediate 12 (1.0g, 2.8mmol), 2-chloronicotinic acid (450mg, 2.86mmol) and NMM (370 $\mu$ l, 3.36mmol) in DMF (30ml). The reaction mixture was stirred overnight at room temperature. The solvent was removed *in vacuo* and the residue partitioned between EtOAc (300ml) and NaHCO<sub>3</sub> solution (300ml). The organic phase was washed with citric acid (10%, 2 x 200ml), NaHCO<sub>3</sub> solution (200ml) and brine (300ml), dried (MgSO<sub>4</sub>) and concentrated *in vacuo* to give the title compound as a yellow oil (1.36g, 98%).  $\delta$ H (CDCl<sub>3</sub>, 300MHz) 9.12 (1H, d,  $\downarrow$  8.0Hz), 8.47 (1H, m), 7.91 (2H, d,  $\downarrow$  8.0Hz), 7.6 (1H, dd), 7.45 (5H, m), 6.95 (2H, d,  $\downarrow$  8.0Hz), 4.67 (1H, m), 4.15 (2H, m), 3.75 (3H, s), 3.31 (1H, m), 3.23 (1H, m) and 1.17 (3H, s);  $m/z$  (ESI) 497 (MH<sup>+</sup>).

**INTERMEDIATE 14****Ethyl 2-(((2-chloro-3-pyridinyl)carbonyl)amino)-3-(4-(carboxyl)phenyl)propanoate**

5 TFA (20ml) was added to a solution of Intermediate 13 (1.36g, 2.75mmol) in toluene (20ml). The reaction mixture was stirred for 30min at room temperature. The white solid obtained was recrystallised (EtOAc/hexane) to give the title compound (1.04g, 100%).  $\delta$ H (CDCl<sub>3</sub>, 300MHz) 12.84 (1H, br s), 9.1 (1H, d,  $\downarrow$  8.0Hz), 8.47 (1H, m), 7.87 (2H, d,  $\downarrow$  8.0Hz), 4.1 (2H, m), 3.2 (1H, m), 3.07 (1H, m) and 1.17 (3H, m);  $m/z$  (ESI) 377 (MH<sup>+</sup>).

**INTERMEDIATE 15****Ethyl 2-(((2-chloro-3-pyridinyl)carbonyl)amino)-3-(4-((2,6-dichloroanilino)carbonyl)phenyl)propanoate**

15 Carbon tetrachloride (5ml) was added to a suspension of Intermediate 14 (1.04g, 2.76mmol) and triphenylphosphine (0.87g, 3.31mmol) in acetonitrile. The reaction mixture was stirred for 2h at room temperature. 2,6-Dichloroaniline (0.89g, 5.52mmol) and NMM (455 $\mu$ l, 4.14mmol) were added and the mixture stirred for a further 48h at room temperature. The solvent was removed *in vacuo* and the residue partitioned between water and EtOAc. The aqueous layer was extracted with EtOAc (x 2) and the combined organic extracts washed with water (x 2) and saturated aqueous NaHCO<sub>3</sub> (x2), dried (Na<sub>2</sub>SO<sub>4</sub>) and concentrated *in vacuo*. Chromatography (SiO<sub>2</sub>; EtOAc/hexane 50:50) gave the title compound (924mg)  $\delta$ H (CDCl<sub>3</sub>, 300MHz) 8.46 (1H, s), 8.05 (1H, d,  $\downarrow$  7.4Hz), 7.89 (2H, d,  $\downarrow$  7.9Hz), 7.69-7.10 (8H, m), 5.11 (1H, dt,  $\downarrow$  6.7, 5.8Hz), 4.25 (2H, q,  $\downarrow$  7.1Hz), 3.42 (1H, dd,  $\downarrow$  13.9, 5.8Hz), 3.31 (1H, dd,  $\downarrow$  13.9, 5.8Hz) and 1.30 (3H, t,  $\downarrow$  7.1Hz).

**INTERMEDIATE 16****Methyl 4-(2(2,6-dichlorophenyl)-2-hydroxyethyl)benzoate**

30 A solution of methyl 4-(bromomethyl)benzoate (2.0g, 8.7mmol) in THF (4.4ml) was added slowly to cut zinc foil (683mg, 10.44mmol) which had been activated with 1,2-dibromoethane (80mg). After 3h of stirring at room temperature 2ml of the solution was transferred to a solution of copper cyanide (396mg, 4.4mmol) and lithium chloride (356mg, 8.4mmol) in THF (4ml) cooled to -78°. This solution was warmed to -20° and then cooled

back to  $-78^{\circ}$ . Boron trifluoride etherate (983  $\mu$ l, 8mmol) was then added followed by 2,6-dichlorobenzaldehyde (0.56g, 3.2mmol) in THF (1ml). The reaction was stirred for 2h and then allowed to warm slowly to room temperature. Water (20ml) was then added and the reaction mixture extracted into EtOAc (3 x 25ml) and the combined organics dried ( $\text{Na}_2\text{SO}_4$ ) and evaporated. Purification by column chromatography ( $\text{SiO}_2$ ; hexane:EtOAc, 4:1) gave the title compound as a colourless oil (863mg, 83%).  $\delta\text{H}$  ( $\text{CDCl}_3$ ) 7.95 (2H, m, ArH), 7.93-7.26 (4H, m, ArH), 7.17-7.12 (1H, m, ArH), 5.73-5.65 (1H, m, CH), 3.91 (3H, s,  $\text{CO}_2\text{Me}$ ), 3.43 (1H, dd,  $\downarrow$  13.5, 8.4Hz,  $\text{CH}_\text{A}\text{H}_\text{B}$ ) and 3.28 (1H, dd,  $\downarrow$  13.5, 6.3Hz,  $\text{CH}_\text{A}\text{H}_\text{B}$ ).  $m/z$  (ESI, 60V) 325 ( $\text{MH}^+$ ).

#### INTERMEDIATE 17

##### Methyl 4-[2-[(tert-butyl(dimethyl)silyl]oxy]-2-(2,6-dichlorophenyl)ethyl]benzoate

To a solution of Intermediate 16 (20g, 6.15mmol) in DCM (10ml) cooled to  $0^{\circ}$  was added 2,4,6-collidine (2.03ml, 15.39mmol). After 15min *tert*-butyldimethylsilyltrifluoromethanesulphonate (2.12ml, 9.23mmol) was added. The reaction mixture was stirred overnight at room temperature then diluted with DCM (100ml) and washed with 1M hydrochloric acid (50ml), water (50ml) and brine (50ml), dried ( $\text{Na}_2\text{SO}_4$ ) and evaporated under reduced pressure. Purification by column chromatography ( $\text{SiO}_2$ ; hexane:EtOAc, 5:1) gave the title compound as a pale pink oil (2.67g, 100%).  $\delta\text{H}$  ( $\text{CDCl}_3$ ) 7.94 (2H, d,  $\downarrow$  6.5Hz, ArH), 7.33, 7.21 (4H, m, ArH), 7.13-7.07 (1H, m, ArH), 5.58 (1H, dd,  $\downarrow$  9.4, 4.6Hz, CH), 3.90 (1H, s,  $\text{CO}_2\text{Me}$ ), 3.46 (1H, dd,  $\downarrow$  13.1, 9.4Hz,  $\text{CH}_\text{A}\text{H}_\text{B}$ ), 3.04 (1H, dd,  $\downarrow$  13.1, 4.6Hz,  $\text{CH}_\text{A}\text{H}_\text{B}$ ), 0.74 (9H, s,  $\text{Si}^t\text{Bu}$ ), -0.31 (3H, s,  $\text{SiMe}$ ) and -0.32 (3H, s,  $\text{SiMe}$ );  $m/z$  (ESI, 60V) 361 ( $\text{MH}^+$ ).

#### INTERMEDIATE 18

##### 4-[2-[(Tert-butyl(dimethyl)silyl]oxy]-2-(2,6-dichlorophenyl)ethyl]benzylalcohol

Lithium aluminium hydride (1M solution in THF, 6.46ml, 6.46mmol) was added to an ice cold solution of Intermediate 17 (2.67g, 6.15mmol) in THF (20ml). The reaction mixture was stirred for 1h then quenched with the addition of water and extracted into DCM (3 x 50ml), dried ( $\text{Na}_2\text{SO}_4$ ) and

evaporated under reduced pressure. Purification by column chromatography (SiO<sub>2</sub>; hexane:EtOAc, 4:1) gave the title compound as a colourless oil (2.g, 87%)  $\delta$ H 7.32-7.06 (7H, m, ArH), 5.56 (1H, dd,  $\downarrow$  9.3, 4.7Hz, CH<sub>2</sub>CH), 4.65 (2H, d,  $\downarrow$  5.9Hz, CH<sub>2</sub>O), 3.39 (1H, dd,  $\downarrow$  13.2, 9.3Hz, CH<sub>A</sub>H<sub>B</sub>), 3.01 (1H, dd,  $\downarrow$  13.2, 4.7Hz, CH<sub>A</sub>H<sub>B</sub>), 0.75 (9H, s, Si<sup>t</sup>Bu), -0.29 (3H, s, SiMe) and -0.31 (3H, s, SiMe);  $m/z$  (ESI, 60V) 433 (MH<sup>+</sup>).

#### **INTERMEDIATE 19**

##### **4-[2-([*Tert*-butyl(dimethyl)silyloxy]-2-(2,6-dichlorophenyl)ethyl]benzylbromide**

A solution of triphenylphosphine (843mg, 3.21mmol) in DCM (2ml) was added to a solution of carbon tetrabromide (1.42g, 3.73mmol) and Intermediate 18 (1.10g, 2.67mmol) in DCM (3ml) and stirred at room temperature for 24h. Ether (100ml) was added and the solid precipitate formed removed by filtration. The filtrate was evaporated under reduced pressure and the residue purified by column chromatography (SiO<sub>2</sub>; 8:1, hexane:EtOAc) to give the title compound (1.20g, 95%).  $\delta$ H (CDCl<sub>3</sub>) 7.32-7.07 (7H, m, ArH), 5.54 (1H, dd,  $\downarrow$  9.5, 4.4Hz, CH<sub>2</sub>CH), 4.49 (2H, s, CH<sub>2</sub>Br), 3.39 (1H, dd,  $\downarrow$  13.2, 9.5Hz, CH<sub>A</sub>H<sub>B</sub>), 2.96 (1H, dd,  $\downarrow$  13.2, 4.4Hz, CH<sub>A</sub>H<sub>B</sub>), 0.73 (9H, s, Si<sup>t</sup>Bu), -0.31 (3H, s, SiMe) and -0.32 (3H, s, SiMe);  $m/z$  (ESI, 60V) 474 (MH<sup>+</sup>).

#### **INTERMEDIATE 20**

##### **4-[2-([*Tert*-butyl(dimethyl)silyloxy]-2-(2,6-dichlorophenyl)ethyl]phenylalanine ethyl ester**

To a solution of ethyl *N*-(diphenylmethylene)glycinate (2.63g, 9.81mmol) in THF (50ml) cooled to -78° was added lithium diisopropylamine (2M in heptane/THF/ethylbenzene, 5.64ml, 11.28mmol). The solution was stirred for 45min. Intermediate 19 (4.20g, 8.92mmol) in THF (20ml) was then added dropwise. The reaction mixture was stirred for 2h at -78° and then warmed to room temperature. EtOAc (100ml) was added and the mixture washed with water (75ml) and brine (75ml), dried (Na<sub>2</sub>SO<sub>4</sub>) and evaporated under reduced pressure. The residue was then taken up in acidic ethanol and stirred for 10min. The volatiles were then removed and the residue partitioned between EtOAc (150ml) and saturated aqueous Na<sub>2</sub>CO<sub>3</sub> (100ml). The aqueous layer was extracted several times with

EtOAc and the combined organics dried ( $\text{Na}_2\text{SO}_4$ ) and evaporated under reduced pressure. The remaining residue was purified by column chromatography ( $\text{SiO}_2$ ; EtOAc) to give the title compound as a colourless oil (3.95g, 95%).  $\delta\text{H}$  ( $\text{CDCl}_3$ ) 7.32-7.05 (7H, m, ArH), 5.52 (1H, dd,  $\downarrow$  9.4, 4.6Hz,  $\text{CHOSi}$ ), 4.18 (2H, q,  $\downarrow$  7.1Hz,  $\text{CO}_2\text{CH}_2\text{CH}_3$ ), 3.68 (1H, dd,  $\downarrow$  7.9, 5.1Hz,  $\text{CHNH}_2$ ), 3.35 (1H, dd,  $\downarrow$  13.2, 9.4Hz,  $\text{CHOSiCH}_2\text{CH}_3$ ), 3.07 (1H, dd,  $\downarrow$  13.5, 5.1Hz,  $\text{CHNH}_2\text{CH}_2\text{CH}_3$ ), 2.95 (1H, dd,  $\downarrow$  13.2, 4.6Hz,  $\text{CHOSiCH}_2\text{CH}_3$ ), 2.95 (1H, dd,  $\downarrow$  13.2, 4.6Hz,  $\text{CHOSiCH}_2\text{CH}_3$ ), 2.82 (1H, dd,  $\downarrow$  13.5, 7.9Hz,  $\text{CHNH}_2\text{CH}_2\text{CH}_3$ ), 1.53 (2H, br s,  $\text{NH}_2$ ), 1.27 (3H, t,  $\downarrow$  7.1Hz,  $\text{CO}_2\text{CH}_2\text{CH}_3$ ), 0.74 (9H, s,  $\text{Si}^t\text{Bu}$ ) and -0.32 (6H, s,  $\text{SiMe}_2$ );  $m/z$  (ESI, 60V) 496 ( $\text{MH}^+$ ).

#### **INTERMEDIATE 21**

##### **4-[2-[(*Tert*-butyl(dimethyl)silyl]oxy]-2-(2,6-dichlorophenyl)ethyl]-(*N*-2-chloronicotinoyl)phenylalanine ethyl ester**

To a solution of Intermediate 20 (1.50g, 3.20mmol) and 2-chloronicotinic acid (504mg, 3.20mmol) in DCM (75ml) at room temperature was added NMM (386 $\mu\text{l}$ , 3.53mmol), EDC (675mg, 3.53mmol) and HOBT (477mg, 3.53mmol). The reaction mixture was stirred overnight at room temperature and then diluted with DCM (50ml) and washed with saturated aqueous  $\text{Na}_2\text{CO}_3$  (50ml), water (50ml) and brine (50ml), dried ( $\text{Na}_2\text{SO}_4$ ) and evaporated under reduced pressure. The residue was purified by column chromatography ( $\text{SiO}_2$ ; EtOAc) to give the title compound as a white solid (1.72g, 85%).  $\delta\text{H}$  ( $\text{CDCl}_3$ ) 8.41-8.38 (1H, m, NH), 8.01-7.96 (1H, m, ArH), 7.29-6.97 (9H, m, ArH), 5.47 (1H, dd,  $\downarrow$  9.2, 4.7Hz,  $\text{CHOSi}$ ), 5.24-4.96 (1H, m,  $\text{CHNH}$ ), 4.19 (2H, qd,  $\downarrow$  7.1, 1.1Hz,  $\text{CO}_2\text{CH}_2\text{CH}_3$ ), 3.34-3.14 (4H, m,  $\text{CH}_2 \times 2$ ), 1.27 (3H, td,  $\downarrow$  7.1, 1.3Hz,  $\text{CO}_2\text{CH}_2\text{CH}_3$ ), 0.69 (s) and 0.66 (s); together (9H,  $\text{Si}^t\text{Bu}$ ) and -0.38 (s) and -0.40 (s); together (6H,  $\text{SiMe}_2$ ); 659 ( $\text{M}^+ + \text{Na}^+$ ).

#### **INTERMEDIATE 22**

##### **(*N*-2-Chloronicotinoyl)-4-[2-(2,6-dichlorophenyl)-2-hydroxyethyl]phenylalanine ethyl ester**

Tetrabutylammonium fluoride (1M in THF, 4.7ml, 4.70mmol) was added to a solution of Intermediate 21 (1.50g, 2.35mmol) in THF (75ml) at room temperature. The reaction mixture was stirred for 2h and then the THF

removed and the residue partitioned between EtOAc and water. The layers were separated and the aqueous layer extracted with EtOAc. The combined organics were dried (Na<sub>2</sub>SO<sub>4</sub>) and evaporated under reduced pressure. Purification of the residue by column chromatography (SiO<sub>2</sub>; MeOH:DCM, 5:95) gave the title compound as a pale brown oil (1.03g, 84%)  $\delta$  H (CDCl<sub>3</sub>) 8.31-8.29 (1H, m, NH), 7.86-7.82 (1H, m, ArH), 7.25-6.96 (9H, m, ArH), 5.54-5.49 (1H, m, CH), 4.94-4.88 (1H, m, CH), 4.11 (2H, qd,  $\downarrow$  7.1, 2.0Hz, CO<sub>2</sub>CH<sub>2</sub>CH<sub>3</sub>), 3.28-3.03 (4H, m, 2 x CH<sub>2</sub>) and 1.19 (3H, td,  $\downarrow$  7.1, 1.1Hz, CO<sub>2</sub>CH<sub>2</sub>CH<sub>3</sub>);  $m/z$  (ESI, 60V) 521 (MH<sup>+</sup>).

#### INTERMEDIATE 23

##### (N-2-Chloronicotinoyl)-[4-[2-(2,6-dichlorophenyl)-2-oxoethyl]]phenylalanine ethyl ester

To a solution of Intermediate 22 (300mg, 0.58mmol) in acetone (20ml) was added Jones' Reagent dropwise until an orange colour persisted. i-Propyl alcohol was added to use up excess reagent and then the solution was basified by the addition of saturated aqueous Na<sub>2</sub>CO<sub>3</sub> solution. The solution was then decanted from the solids and the acetone removed *in vacuo*. The remaining aqueous solution was then extracted with ether (x 2) and the combined organics dried (Na<sub>2</sub>SO<sub>4</sub>) and evaporated under reduced pressure. The residue was purified by column chromatography (SiO<sub>2</sub>; hexane:EtOAc, 3:2) to give the title compound as a colourless oil (200mg, 67%).  $\delta$  H (CDCl<sub>3</sub>) 8.40 (1H, dd,  $\downarrow$  4.8, 2.0Hz), 7.94 (1H, dd,  $\downarrow$  7.7, 2.0Hz), 7.38-7.11 (8H, m, ArH), 7.02 (1H, d,  $\downarrow$  7.5Hz, ArH), 4.19 (2H, q,  $\downarrow$  7.2Hz, CO<sub>2</sub>CH<sub>2</sub>CH<sub>3</sub>), 4.09 (2H, s, CH<sub>2</sub>CO), 3.28 (1H, dd,  $\downarrow$  14.0, 5.9Hz, CH<sub>A</sub>H<sub>B</sub>), 3.18 (1H, dd,  $\downarrow$  14.0, 6.1Hz, CH<sub>A</sub>H<sub>B</sub>) and 1.26 (5H, t,  $\downarrow$  7.2Hz, CO<sub>2</sub>CH<sub>2</sub>CH<sub>3</sub>);  $m/z$  (ESI, 60V) 519 (MH<sup>+</sup>).

#### INTERMEDIATE 24

##### Methyl 4-[(E)-2-(2,6-dichlorophenyl)ethenyl]benzoate

A solution of Intermediate 16 (2.0g, 6.15mmol) in toluene (25ml) containing p-toluenesulphonic acid (100mg) was heated to reflux in a Dean-Stark apparatus for 4h. Toluene was then removed under reduced pressure and the residue purified by column chromatography (SiO<sub>2</sub>; hexane:EtOAc, 5:1) to give the title compound as an off white solid (1.64g, 87%);  $\delta$  H (CDCl<sub>3</sub>) 8.05 (2H, d,  $\downarrow$  8.3Hz), 7.60 (2H, d,  $\downarrow$  8.3Hz, ArH), 7.36 (2H, d,  $\downarrow$  8.0Hz),

7.21 (2H, d,  $J$  1.7Hz), 7.15-7.10 (1H, m), and 3.93 (3H, s,  $\text{CO}_2\text{CH}_3$ );  $m/z$  (ESI, 60V) 329 ( $\text{MH}^+$ ).

**INTERMEDIATE 25**

5 4-[(E)-2-(2,6-Dichlorophenyl)ethenyl]benzyl alcohol

To an ice cold solution of Intermediate 24 (1.56g, 5.08mmol) in THF (20ml) was added lithium aluminium hydride (1M in THF, 5.34ml, 5.34mmol). The reaction mixture was stirred for 30min and then quenched by the addition of water (10ml). The resulting biphasic solution was filtered through Celite® and then extracted with DCM (2 x 50ml). The combined organics were dried (Na<sub>2</sub>SO<sub>4</sub>) and evaporated under reduced pressure to give the title compound as a colourless oil which solidified on standing (1.5g, 99%).  $\delta$ H (CDCl<sub>3</sub>) 7.55 (2H, d,  $\downarrow$  8.2Hz), 7.42-7.33 (4H, m), 7.20-7.08 (3H, m) and 4.75 (2H, d,  $\downarrow$  5.3Hz, CH<sub>2</sub>); m/z (ESI, 60V) 301 (M<sup>++</sup> Na<sup>+</sup>).

**INTERMEDIATE 26**

**4-[(E)-2-(2,6-Dichlorophenyl)ethenyl]benzyl bromide**

A solution of triphenylphosphine (1.58g, 6.04mmol) in DCM (10ml) was added to a solution of Intermediate 25 (1.40g, 5.03mmol) and carbon tetrabromide (2.33g, 7.04mmol) in DCM (10ml). The resulting solution was stirred for 1h and then diluted with ether (150ml) and the resulting solid removed by filtration. The filtrate was then evaporated under reduced pressure and the resulting residue purified by column chromatography (SiO<sub>2</sub>; hexane: EtOAc, 6:1) to give the title compound as a colourless oil (1g, 49%).  $\delta$ H (CDCl<sub>3</sub>) 7.55 (2H, d,  $J$  8.3Hz), 7.44-7.32 (3H, m), 7.23-7.09 (4H, m) and 4.53 (2H, s, CH<sub>2</sub>);  $m/z$  (ESI, 60V) 342 (MH<sup>+</sup>)

## INTERMEDIATE 27

**4-[(E)-2-(2,6-Dichlorophenyl)ethenyl]phenylalanine ethyl ester**

30 To a solution of ethyl *N*-(diphenylmethylene)glycinate (860mg, 3.21mmol) in THF (20ml) cooled to -78° was added lithium diisopropylamine (2M in heptane/THF/ethylbenzene, 1.68ml, 3.36mmol). The solution was stirred for 45min. Intermediate 26 (10g, 2.92mmol) in THF (2ml) was added and the resulting reaction mixture stirred for 2h at -78° and then warmed to  
35 room temperature. EtOAc (100ml) was added and the mixture washed with water (75ml) and brine (75ml), dried (Na<sub>2</sub>SO<sub>4</sub>) and evaporated under

reduced pressure. The residue was then taken up in acidic ethanol and stirred for 1min. The volatiles were then removed and the residue partitioned between EtOAc (150ml) and saturated aqueous Na<sub>2</sub>CO<sub>3</sub> (100ml). The aqueous layer was extracted several times with EtOAc and the combined organics dried (Na<sub>2</sub>SO<sub>4</sub>) and evaporated under reduced pressure. The remaining residue was purified by column chromatography (SiO<sub>2</sub>; EtOAc) to give the title compound as a colourless oil (860mg, 81%).

$\delta$ H (CDCl<sub>3</sub>) 7.47 (2H, d,  $\downarrow$  8.1Hz), 7.32 (2H, d,  $\downarrow$  8.1Hz), 7.21 (2H, d,  $\downarrow$  8.2Hz), 7.19-6.98 (3H, m), 4.18 (2H, q,  $\downarrow$  7.1Hz, CO<sub>2</sub>CH<sub>2</sub>CH<sub>3</sub>), 3.71 (1H, dd,  $\downarrow$  7.8, 5.3Hz, CH), 3.10 (1H, dd,  $\downarrow$  13.5, 5.3Hz, CH<sub>A</sub>H<sub>B</sub>), 2.88 (1H, dd,  $\downarrow$  13.5, 7.8Hz, CH<sub>A</sub>H<sub>B</sub>) and 1.26 (3H, t,  $\downarrow$  7.1Hz, CO<sub>2</sub>CH<sub>2</sub>CH<sub>3</sub>);  $m/z$  (ESI, 60V) 364 (MH<sup>+</sup>)

#### **INTERMEDIATE 28**

##### **15 (N-2-Chloronicotinoyl)-4-[(E)-2-(2,6-dichlorophenyl)ethenyl]phenylalanine ethyl ester**

To a solution of Intermediate 27 (860mg, 2.36mmol) and 2-chloronicotinic acid (372mg, 2.36mmol) in DCM (25ml) was added NMM (285 $\mu$ l, 2.60mmol), EDC (498mg, 2.60mmol) and HOBT (352mg, 2.60mmol). The resulting solution was stirred for 3h and then diluted with DCM (50ml) and brine (50ml), dried (Na<sub>2</sub>SO<sub>4</sub>) and evaporated under reduced pressure. The residue was purified by column chromatography (SiO<sub>2</sub>; EtOAc) to give the title compound as a pale yellow oil (1.1g, 93%).

$\delta$ H (CDCl<sub>3</sub>) 8.43 (1H, dd,  $\downarrow$  4.8, 2.0Hz), 8.02 (1H, dd,  $\downarrow$  7.6, 2.0Hz), 7.46 (2H, d,  $\downarrow$  8.2Hz), 7.34-6.92 (9H, m), 5.10-5.03 (1H, m, CH), 4.24 (2H, q,  $\downarrow$  7.1Hz, CO<sub>2</sub>CH<sub>2</sub>CH<sub>3</sub>), 3.33 (1H, dd,  $\downarrow$  13.9, 5.8Hz, CH<sub>A</sub>H<sub>B</sub>), 3.24 (1H, dd,  $\downarrow$  13.9, 5.8Hz, CH<sub>A</sub>H<sub>B</sub>) and 1.30 (3H, t,  $\downarrow$  7.1Hz, CO<sub>2</sub>CH<sub>2</sub>CH<sub>3</sub>);  $m/z$  (ESI, 60V) 503.5 (MH<sup>+</sup>).

#### **INTERMEDIATE 29**

##### **30 N-(2-Chloronicotinoyl)-L-tyrosine methyl ester**

EDC.HCl (2.11g, 11mmol) was added to a mixture of L-tyrosine methyl ester hydrochloride (2.32g, 10mmol), 2-chloronicotinic acid (1.58g, 10mmol), HOBT (1.49g, 11mmol) and NMM (2.31ml, 21mmol) in DMF (50ml). The mixture was stirred overnight at room temperature. The solvent was removed *in vacuo*. the residue dissolved in EtOAc (300ml) and washed with dilute HCl (100ml), saturated aqueous NaHCO<sub>3</sub>



(100ml), water (3 x 100ml) and brine (50ml), dried ( $\text{Na}_2\text{SO}_4$ ) and solvent removed *in vacuo* to give the title compound as a yellow gum (3.27g, 98%).  $\delta\text{H}$  ( $\text{DMSO}-d_6$ , 300MHz) 9.21 (1H, s, OH), 9.03 (1H, d,  $\downarrow$  7.9Hz, CONH), 8.45 (1H, dd,  $\downarrow$  4.8, 1.9Hz, pyrH), 7.67 (1H, dd,  $\downarrow$  7.4, 1.9Hz, pyrH), 7.47 (1H, dd,  $\downarrow$  7.5, 4.8Hz, pyrH), 7.05 (2H, d,  $\downarrow$  8.5Hz, ArH), 6.67 (2H, d,  $\downarrow$  8.5Hz, ArH), 4.58 (1H, ddd,  $\downarrow$  9.6, 7.9, 5.4Hz,  $\text{CH}_\alpha$ ), 3.65 (3H, s,  $\text{CO}_2\text{Me}$ ), 3.01 (1H, dd,  $\downarrow$  13.9, 5.4Hz,  $\text{CH}_\text{A}\text{H}_\text{B}\text{Ar}$ ) and 2.85 (1H, dd,  $\downarrow$  13.9, 9.6Hz,  $\text{CH}_\text{A}\text{H}_\text{B}\text{Ar}$ ),  $m/z$  (ESI, 60V) 335 ( $\text{MH}^+$ ).

#### 10 INTERMEDIATE 30

##### *N*-(2-Chloronicotinoyl)-*O*-(2,6-dichlorobenzoyl)-*L*-tyrosine methyl ester

A solution of Intermediate 29 (919mg, 2.75mmol) in DMF (10ml) was added to a suspension of sodium hydride (60% in mineral oil, 3.03mmol, 121mg) in DMF (20ml) at  $0^\circ$ . After 15min, 2,6-dichlorobenzoyl chloride (414 $\mu\text{l}$ , 2.89mmol) was added and the mixture stirred for 2h at room temperature. Water (~5ml) was added and the solvent removed *in vacuo*. The residue was dissolved in EtOAc (150ml), washed with water (3 x 50ml) and brine (50ml), dried ( $\text{Na}_2\text{SO}_4$ ) and evaporated *in vacuo*. Chromatography ( $\text{SiO}_2$ ; DCM/MeOH, 98:2) gave the title compound as a white foam (1.10g, 79%).  $\delta\text{H}$  ( $\text{DMSO}-d_6$ , 300MHz) 9.12 (1H, d,  $\downarrow$  7.6Hz, CONH), 8.45 (1H, dd,  $\downarrow$  4.8, 2.0Hz, pyrH), 7.69-7.58 (4H, m, pyrH +  $\text{Cl}_2\text{ArH}_3$ ), 7.46 (1H, dd,  $\downarrow$  7.5, 4.9Hz, pyrH), 7.43 (2H, d,  $\downarrow$  8.4Hz,  $\text{ArH}_2$ ), 7.23 (2H, d,  $\downarrow$  8.5Hz,  $\text{ArH}_2$ ), 4.73 (1H, m,  $\text{CH}_\alpha$ ), 3.68 (3H, s,  $\text{CO}_2\text{Me}$ ), 3.22 (1H, dd,  $\downarrow$  13.9, 5.2Hz,  $\text{CH}_\text{A}\text{H}_\text{B}\text{Ar}$ ) and 3.02 (1H, dd,  $\downarrow$  13.9, 10.1Hz,  $\text{CH}_\text{A}\text{H}_\text{B}\text{Ar}$ );  $m/z$  (ESI, 60V) 507 ( $\text{MH}^+$ ).

#### INTERMEDIATE 31

##### *N*-(2-Chloronicotinoyl)-*N*-methyl-*N'*-(3,5-dichloro-4-picolyl)-*L*-4-aminophenylalanine methyl ester

2-Chloronicotinoyl chloride (132mg, 0.75mmol) was added to *N*-methyl-*N'*-(3,5-dichloroisonicotinyl)-*L*-4-aminophenylalanine methyl ester [prepared from *N*-Boc-*N'*-phthaloyl-4-amino-*L*-phenylalanine methyl ester and methyl iodide, followed by treatment with hydrazine monohydrate and reaction with 3,5-dichloroisonicotinyl chloride with subsequent removal of the Boc group] and NMM (165 $\mu\text{l}$ , 1.5mmol) in DCM (10ml). The mixture was stirred for 1h

at room temperature then diluted with DCM (100ml) and washed with dilute HCl (30ml), dried (Na<sub>2</sub>SO<sub>4</sub>) and evaporated *in vacuo*. Chromatography (SiO<sub>2</sub>; EtOAc/hexane, 10:90) gave the title compound as a colourless gum (380mg, 97%).  $\delta$ H (DMSO-d<sub>6</sub>, 300MHz, 405K) 10.35 (1H, br s, CONH), 8.67 (2H, s, Cl<sub>2</sub>pyrH), 8.42 (1H, t,  $\Delta$  3.4Hz, ClpyrH), 7.55 (2H, br d,  $\Delta$  7.2Hz, ArH), 7.45-7.15 (4H, v br m, ArH + ClpyrH), 5.3 (1H, v br s, CH $\alpha$ ), 3.74 (3H, s, CO<sub>2</sub>Me), 3.4-3.3 (1H, br m, CH<sub>A</sub>H<sub>B</sub>Ar), 3.16 (1H, dd,  $\Delta$  14.4, 9.6Hz, CH<sub>A</sub>H<sub>B</sub>Ar) and 2.73 (3H, br s, NMe);  $m/z$  (ESI, 60V) 521 (MH<sup>+</sup>).

### INTERMEDIATE 32

#### [(S-2,5-dimethoxyphenyl)sulphonyl]nicotinoyl-O-(2,6-dichlorobenzyl)-L-tyrosine methyl ester

A solution of Intermediate 2 (0.79g, 1.26mmol) in DCM (50ml) was treated with 3-chloroperoxybenzoic acid (2.17g, 12.6mmol) and stored at 4° for 48h. The reaction was partitioned between DCM (20ml) and NaHCO<sub>3</sub> solution (20ml). The aqueous layer was extracted with DCM (25ml) and the combined organic layers washed with 10% aqueous Na<sub>2</sub>SO<sub>3</sub> (50ml), dried (MgSO<sub>4</sub>) and the solvent removed *in vacuo* to give a yellow foam that was purified by chromatography (SiO<sub>2</sub>; EtOAc/hexane 3:1) to give the title compound as a white foam, (0.50g, 60%).  $\delta$ H (CDCl<sub>3</sub>) 8.58 (1H, dd,  $\Delta$  4.7, 1.7Hz, pyr-H), 7.95 (1H, dd,  $\Delta$  7.8, 1.7Hz, pyr-H), 7.70 (1H, d,  $\Delta$  3.2Hz, Ar-H), 7.50 (1H, dd,  $\Delta$  7.8, 4.7Hz, pyr-H), 7.36-7.11 (6H, m, Ar-H), 7.01 (1H, d,  $\Delta$  7.6Hz, NH), 6.92-6.84 (3H, m, Ar-H), 5.21 (2H, s, CH<sub>2</sub>O), 5.10 (1H, m, CH $\alpha$ ), 3.85 (3H, s, OMe), 3.74 (3H, s, OMe), 3.49 (3H, s, CO<sub>2</sub>Me) and 3.27 (2H, m, CHCH<sub>2</sub>Ar).  $m/z$  (ESI, 60V) 659 (MH<sup>+</sup>).

### EXAMPLE 1

#### 2-Chloronicotinoyl-O-(2,6-dichlorobenzyl)-L-tyrosine

A solution of Intermediate 1 (0.20g, 0.41mmol) in THF (5ml) and water (5ml) was treated with lithium hydroxide monohydrate (25ml, 0.61mmol, 1.5 equiv.) and stirred at room temperature for 1.5h. The reaction was acidified to pH1 with 10% hydrochloric acid to give a white precipitate which was isolated by filtration, washed with water (5ml) and diethyl ether (5ml) and dried under vacuum to give the title compound as a white

- powder (0.14g, 72%).  $\delta$ H (DMSO- $d_6$ ) 8.93 (1H, d,  $\downarrow$  8.1Hz), 8.45 (1H, dd,  $\downarrow$  2.0, 4.8Hz), 7.66 (1H, dd,  $\downarrow$  2.0, 7.5Hz), 7.56 (2H, m), 7.47 (2H, m), 7.24 (2H, d,  $\downarrow$  8.6Hz), 6.99 (2H, d,  $\downarrow$  8.6Hz), 5.20 (2H, s), 4.60 (1H, d), 3.13 (1H, ABX,  $\downarrow$  4.7, 13.9Hz) and 2.90 (1H, ABX,  $\downarrow$  10.0, 13.9Hz).  $m/z$  (ES+, 60V) 479, 481 (MH+).

## EXAMPLE 2

### 2-Thio(S-2,5-dimethoxyphenyl)nicotinoyl-O-(2,6-dichlorobenzyl)-L-tyrosine hydrochloride

- 10 A solution of Intermediate 2 (0.35g, 0.56mmol) in THF (15ml) and water (7.5ml) was treated with lithium hydroxide monohydrate (28mg, 0.67mmol) and stirred at room temperature for 16h. The reaction was acidified to pH1 with 10% hydrochloric acid, extracted with DCM (2 x 30ml), and the combined organic layers were dried over  $MgSO_4$ , the solvent removed *in vacuo* to give a gummy residue, which was purified by chromatography (SiO<sub>2</sub>; 7.5% MeOH/DCM) to give a gum which was dissolved in acetonitrile (20ml) and water (20ml) and lyophilised to give the title compound as a white powder (0.26g, 71%).  $\delta$ H (DMSO- $d_6$ ) 8.71 (1H, m), 8.29 (1H, dd,  $\downarrow$  1.7, 4.8Hz), 7.75 (1H, d,  $\downarrow$  6.0Hz), 7.56-6.94 (11H, m), 5.16 (2H, s), 4.56 (1H, m), 3.68 (3H, s), 3.60 (3H, s) and 3.21-1.96 (2H, m).  $m/z$  (ES+ 60V) 613, 615 (MH+).

## EXAMPLE 3

### a) 2-Thio(S-4-picolinyl)nicotinoyl-O-(2,6-dichlorobenzyl)-L-tyrosine

- 25 A solution of Intermediate 4 (0.30g, 0.52mmol) in THF (7.5ml) and water (5ml) was treated with lithium hydroxide monohydrate (33mg, 0.7mmol) and stirred at room temperature for 16h. The pH was adjusted to 6.5-7 with 10% hydrochloric acid to give a yellow precipitate which was isolated by filtration, washed with water and dried *in vacuo* to give the title compound as a yellow powder (0.28g, 95%).  $\delta$ H (DMSO- $d_6$ ) 8.83 (1H, d,  $\downarrow$  8.0Hz), 8.50 (3H, m), 7.75 (1H, d,  $\downarrow$  7.6Hz), 7.56-7.20 (8H, m), 6.95 (2H, d,  $\downarrow$  8.5Hz), 5.17 (2H, s), 4.54 (1H, m), 4.37 (2H, s), 3.12 (1H, m) and 2.97 (1H, m).  $m/z$  (ES+, 60V) 568, 570 (MH+).

- 35 The following compounds were prepared in a similar manner by hydrolysis of the corresponding methyl ester. Each ester starting material was obtained either by alkylation of Intermediate 3 or alternative

mercaptopyridine using the reagents shown using a similar procedure to that described for Intermediate 4:

- b) **2-Thio-S-benzyl-nicotinoyl-(O-2,6-dichlorobenzyl)-L-tyrosine**  
 5 ester from Intermediate 3 and benzyl chloride.  $\delta$ H (DMSO- $d_6$ ) 8.78 (1H, d,  $\downarrow$  8.1Hz, pyr-H), 8.54 (1H, dd,  $\downarrow$  4.8, 1.1Hz, pyr-H), 7.69 (1H, dd,  $\downarrow$  7.7, 1.7Hz, pyr-H), 7.68-7.17 (11H, m, Ar-H), 6.94 (2H, d,  $\downarrow$  8.6, Ar-H), 5.16 (2H, s, CH<sub>2</sub>O), 4.53 (1H, m, CH $\alpha$ ), 4.34 (2H, s, CH<sub>2</sub>Ar), 3.10 (1H, dd,  $\downarrow$  13.9, 4.6Hz, CHCH<sub>A</sub>H<sub>B</sub>Ar), 2.94 (1H, dd,  $\downarrow$  13.9, 10.2Hz, CHCH<sub>A</sub>H<sub>B</sub>Ar);  
 10  $m/z$  (ESI, 60V) 567 (MH<sup>+</sup>).
- c) **2-Thio-(S-4-Methylphenyl)-nicotinoyl-(O-2,6-dichlorobenzyl)-L-tyrosine**  
 ester from Intermediate 3 and 4-methylbenzyl chloride.  $\delta$ H (DMSO- $d_6$ )  
 15 8.87 (1H, d,  $\downarrow$  8.1Hz, pyr-H), 8.31 (1H, dd,  $\downarrow$  4.7, 1.8Hz, pyr-H), 7.70 (1H, dd,  $\downarrow$  7.6, 1.7Hz, pyr-H), 7.54 (2H, m, Ar-H), 7.45 (1H, m, Ar-H), 7.32-7.17 (7H, m, Ar-H), 6.98 (2H, d,  $\downarrow$  8.6Hz, Ar-H), 5.17 (2H, s, CH<sub>2</sub>O), 4.58 (1H, m, CH $\alpha$ ), 3.16 (1H, dd,  $\downarrow$  14.6, 14.0Hz, CHCH<sub>A</sub>H<sub>B</sub>Ar), 2.98 (1H, dd,  $\downarrow$  14.0, 10.1Hz, CHCH<sub>A</sub>H<sub>B</sub>Ar) and 2.32 (3H, s, Me);  $m/z$  (ESI, 60V) 569 (MH<sup>+</sup>).  
 20
- d) **2-Thio-S-(3-picolyl)-nicotinoyl-(O-2,6-dichlorobenzyl)-L-tyrosine**  
 ester from Intermediate 3 and 3-picolyl chloride.  $\delta$ H (DMSO- $d_6$ ) 9.04-  
 8.54 (5H, m, Ar-H), 7.95-7.84 (2H, m, Ar-H), 7.56-7.43 (3H, m, Ar-H), 6.96  
 (2H, d,  $\downarrow$  8.3Hz, Ar-H), 5.17 (2H, s, CH<sub>2</sub>O), 4.55 (1H, m, CH $\alpha$ ), 4.48 (2H,  
 25 s, CH<sub>2</sub>pyr), 3.05 (2H, m, CHCH<sub>2</sub>Ar);  $m/z$  (ESI, 60V) 569 (MH<sup>+</sup>).
- e) **N-[2-thio(S-3-picolinyl)nicotinoyl]-O-2,6-dichlorobenzyl-L-tyrosine**  
 ester from Intermediate 3 and 3-picolyl chloride using DBU as base.  $\delta$ H  
 30 (DMSO- $d_6$ ) 8.8 (1H, br d), 8.57 (1H, m), 8.50 (1H, m), 7.72 (2H, m), 7.55 (2H, d), 7.5-7.4 (2H, m), 7.3-7.15 (4H, m), 6.95 (2H, d), 5.15 (2H, s), 4.55-4.45 (3H, m), 3.2-3.1 (1H, m), 3.0-2.9 (1H, m);  $m/z$  (ESI, 60V) 568 (MH<sup>+</sup>).
- f) **N-2-Thio(S-4-butanoate)nicotinoyl-(O-2,6-dichlorobenzyl)-L-tyrosine**  
 35

ester from Intermediate 3 and methyl-4-chlorobutyrate using  $K_2CO_3$  as base.  $\delta H$  (DMSO- $d_6$ ) 8.76 (1H, d,  $\downarrow$  8.1Hz), 8.49 (1H, dd,  $\downarrow$  4.8, 1.7Hz), 7.61 (1H, dd,  $\downarrow$  7.7, 1.7Hz), 7.55 (2H, d,  $\downarrow$  8.9Hz), 7.45 (1H, m), 7.23 (2H, d,  $\downarrow$  8.5Hz), 7.2 (1H, m), 6.96 (2H, d,  $\downarrow$  8.5Hz), 5.18 (2H, s), 4.53 (1H, m), 3.17-2.87 (4H, m), 2.40-2.30 (2H, t,  $\downarrow$  7.3Hz), 1.83 (2H, m);  $m/z$  (ESI, 60V) 563 (MH<sup>+</sup>).

**g) (N'-3,5-Dichloroisonicotinoyl)-N-((3-pyridinylmethyl)thio)isonicotinoyl)-L-4-aminophenylalanine**

ester from 2-mercaptoisonicotinoyl-(N'-3,5-dichloroisonicotinoyl)-L-4-aminophenylalanine and 3-picolyl chloride.  $\delta H$  (DMSO- $d_6$ ) 10.86 (1H, s,  $CO_2H$ ), 8.92-8.75 (3H, m, ArH), 8.68-8.52 (2H, m), 8.39 (1H, br s), 7.75 (2H, t,  $\downarrow$  7.5Hz), 7.56 (2H, d,  $\downarrow$  6.3Hz), 7.39-7.18 (4H, m), 4.60-4.47 (1H, m, CH) and 3.25-2.92 (2H, m,  $CH_2$ );  $m/z$  (ESI, 60V) 582 (MH<sup>+</sup>).

**EXAMPLE 4**

**a) 2-Thio(S-2,5-dimethoxyphenyl)nicotinoyl-(N-2,6-dichlorobenzoyl)-L-4-aminophenylalanine**

A solution of Intermediate 6 (0.44g, 0.69mmol) in THF (7.5ml) and water (5ml) was treated with lithium hydroxide monohydrate (43mg, 1.0mmol) and stirred at room temperature for 16h, then acidified to pH1 with 10% hydrochloric acid. The mixture was extracted with DCM (2 x 30ml) and the solvent evaporated *in vacuo* to give an off-white solid that was triturated with boiling MeOH to give the title compound as a white solid (210mg, 49%).  $\delta H$  (DMSO- $d_6$ ) 11.18 (1H, br s,  $CO_2H$ ), 10.66 (1H, s, NH), 8.89 (1H, d,  $\downarrow$  8.0Hz), 8.30 (1H, dd,  $\downarrow$  1.6, 4.7Hz), 7.77 (1H, dd,  $\downarrow$  1.6, 7.6Hz), 7.62-7.45 (5H, m), 7.32 (2H, d,  $\downarrow$  8.5Hz), 7.17 (1H, dd,  $\downarrow$  4.8, 7.6Hz), 6.96 (3H, m), 4.62 (1H, m), 3.70 (3H, s), 3.60 (3H, s) and 3.20-3.00 (2H, m).  $m/z$  (ES<sup>+</sup>) 626,628 (MH<sup>+</sup>).

The following compounds were prepared in a similar manner to the compound of Example 4a) by hydrolysis of the corresponding methyl ester. Each ester was obtained by coupling the starting materials shown according to the method described for Intermediate 6:

**a) 2-Thio(S-2,5-dimethoxyphenyl)nicotinoyl-(N-2,6-**

**dichlorobenzoyl)-L-4-aminophenylalanine**

from (N-2,6-dichlorobenzoyl)-L-4-aminophenylalanine methyl ester and 3,5-dichloropyridyl-4-carbonyl chloride.  $\delta$ H (DMSO- $d_6$ ) 8.64 (2H, s), 7.60-7.46 (5H, m), 7.24 (2H, d,  $\downarrow$  8.5Hz), 4.78-4.65 (1H, m), 3.22-2.85 (2H, m);  $\delta$ H  $m/z$  (ESI, 60V) 528 (MH<sup>+</sup>).

c) **(N'-2,4-Dimethylnicotinoyl)-(N-2,6-dichlorobenzoyl)-L-4-aminophenylalanine**

from (N-2,6-dichlorobenzoyl)-L-4-aminophenylalanine methyl ester and 2,4-dimethylpyridyl-4-carbonyl chloride.  $\delta$ H (DMSO- $d_6$ ) 8.39 (1H, br d), 8.23 (1H, d,  $\downarrow$  5.0Hz), 7.59-7.45 (6H, m), 7.26 (2H, d,  $\downarrow$  8.4Hz), 7.01 (1H, d,  $\downarrow$  5.1Hz), 4.65-4.52 (1H, m), 3.26-3.18 (1H, m), 2.95-2.84 (1H, m), 2.17 (3H, s), 2.01 (3H, s);  $m/z$  (ESI, 60V) 486 (MH<sup>+</sup>).

d) **N-(2,6-Dichloroisonicotinoyl)-O-(2,6-dichlorobenzyl)-L-tyrosine**  
from O-(2,6-dichlorobenzyl)-L-tyrosine methyl ester hydrochloride and 2,6-dichloropyridyl-4-carbonyl chloride.  $\delta$ H (DMSO- $d_6$ ) 9.2 (1H, d), 7.81 (2H, s), 7.5-7.3 (3H, m), 7.21 (2H, d,  $\downarrow$  8.5Hz), 6.93 (2H, d,  $\downarrow$  8.5Hz), 5.15 (2H, s), 4.65 (1H, m), 3.28-3.15 (1H, m), 3.05-2.95 (1H, m);  $m/z$  (ESI, 60V) 513 (MH<sup>+</sup>).

e) **N'-(2-nicotinoyl)-O-(2,6-dichlorobenzyl)-L-tyrosine**  
from O-(2,6-dichlorobenzyl)-L-tyrosine methyl ester hydrochloride and nicotinoyl chloride using triethylamine as base.  $\delta$ H (DMSO- $d_6$ ) 9.0-8.85 (2H, m), 8.7 (1H, m), 8.14 (1H, m), 7.55-7.41 (4H, m), 7.24 (2H, d,  $\downarrow$  8.6Hz), 6.95 (2H, d,  $\downarrow$  8.6Hz), 5.16 (2H, s), 4.59 (1H, m), 3.17-3.12 (1H, m), 3.04-2.96 (1H, m);  $m/z$  (ESI, 60V) 445 (MH<sup>+</sup>).

f) **N-(3,5-Dichloro-4-picolyl)-N'-(3,5-dichloro-4-picolyl)-L-4-aminophenylalanine**  
from (N-3,5-dichloro-4-picolyl)-L-4-aminophenylalanine methyl ester and 3,5-dichloropyridyl-4-carbonyl chloride.  $\delta$ H (DMSO- $d_6$ , 300MHz) 9.26 (1H, d,  $\downarrow$  8.3Hz), 8.79 (2H, s), 8.65 (2H, s), 7.57 (2H, d,  $\downarrow$  8.4Hz), 7.30 (2H, d,  $\downarrow$  8.4Hz), 4.70 (1H, m), 3.15 (1H, dd,  $\downarrow$  14.1, 5.2Hz) and 2.93 (1H, dd,  $\downarrow$  14.0, 9.3Hz);  $m/z$  (ESI, 160V) 527 (MH<sup>+</sup>).

g) **N-(2-Chloronicotinoyl)-N'-(3,5-dichloro-4-picolyl)-L-4-amino-phenylalanine**

from (N-3,5-dichloro-4-picolyl)-L-4-aminophenylalanine methyl ester and 2-chloro-nicotinoyl chloride.  $\delta$ H (DMSO- $d_6$ , 300MHz) 12.85 (1H, br s), 10.88 (1H, s), 8.97 (1H, d,  $\downarrow$  8.1Hz), 8.79 (2H, s), 8.46 (1H, dd,  $\downarrow$  4.8, 1.8Hz), 7.70 (1H, dd,  $\downarrow$  7.5, 1.8Hz), 7.59 (2H, d,  $\downarrow$  8.4Hz), 7.48 (1H, dd,  $\downarrow$  7.5, 4.8Hz), 7.30 (2H, d,  $\downarrow$  8.4Hz), 4.63 (1H, m), 3.16 (1H, dd,  $\downarrow$  13.9, 4.7Hz) and 2.95 (1H, dd,  $\downarrow$  13.8, 9.8Hz)  $m/z$  (ESI, 160V) 493 (MH<sup>+</sup>).

10 **EXAMPLE 5**

a) **O-(2,6-dichlorobenzyl)-N-(4-acetyl-1,2,5-trimethyl-3-pyrrolyl)-L-tyrosine**

Intermediate 7 (360mg, 0.68mmol) was treated with LiOH.H<sub>2</sub>O (34mg, 0.81mmol) in dioxane (6ml), water (6ml) and MeOH (4ml) at room temperature for 2h. The solvent was removed *in vacuo* and the obtained residue taken up in water. The pH was made acidic by addition of a few drops of acetic acid and the obtained precipitate filtered off with water washing affording the title compound as a white amorphous powder (245mg, 70%).  $\delta$ H ( $d_6$ -DMSO) 8.37 (1H, d,  $\downarrow$  8.2Hz, NH), 7.57-7.43 (3H, m's, aryl-H), 7.23 (2H, d,  $\downarrow$  8.6Hz, aryl-H), 6.95 (2H, d,  $\downarrow$  87.6Hz, aryl-H), 5.18 (2H, br s, CH<sub>2</sub>-O), 4.62 (1H, m,  $\alpha$ tyr-H), 3.33 (3H, s, MeN), 3.13 (1H, dd,  $\downarrow$  4.1, 13.8Hz, CH<sub>A</sub>H<sub>B</sub>Ar), 2.32 (3H, s, MeCO), 2.01 (3H, s, pyrrole-Me) and 1.92 (3H, s, pyrrole-Me).  $m/z$  (ES<sup>+</sup>, 60V), 517 (MH<sup>+</sup>, 100), 519 (MH<sup>+</sup>, 70).

25 The following compounds were prepared in a similar manner to the compound of Example 5a) by hydrolysis of the corresponding methyl ester. Each ester was obtained by coupling the starting materials shown according to the method described for Intermediate 7:

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b) **O-(2,6-dichlorobenzyl)-N-(4-acetyl-3,5-dimethyl-2-pyrrolyl)-L-tyrosine**

from O-(2,6-dichlorobenzyl)-L-tyrosine methyl ester hydrochloride and 3,5-dimethyl-4-acetylpyrrole-2-carboxylic acid. Freeze drying afforded the title compound as a light cream amorphous solid (550mg).  $\delta$ H (DMSO- $d_6$ ) 11.56, (1H, s), 7.61 (1H, d,  $\downarrow$  7.8Hz), 7.51 (2H, d,  $\downarrow$  8.0Hz), 7.41 (1H, t,  $\downarrow$

35

8.0Hz), 7.21 (2H, d,  $\downarrow$  8.5Hz), 6.96 (2H, d,  $\downarrow$  8.5Hz), 5.16 (2H, s), 4.69-4.55 (1H, m), 3.12 (1H, dd,  $\downarrow$  13.7, 4.7Hz), 2.99 (1H, dd,  $\downarrow$  13.7, 9.1Hz), 2.43 (3H, s), 2.38 (3H, s), 2.31 (3H, s);  $m/z$  (ESI, 60V) 503 (MH<sup>+</sup>).

5 c) **O-(2,6-dichlorobenzyl)-N-(4-acetyl-2,5-dimethyl-3-pyrrolyl)-L-tyrosine**

from O-(2,6-dichlorobenzyl)-L-tyrosine methyl ester hydrochloride and 2,5-dimethyl-4-acetyl-pyrrole-3-carboxylic acid. Freeze drying afforded the title compound as a white amorphous solid (203mg).  $\delta$ H (DMSO-d<sub>6</sub>) 11.2 (1H, s), 8.83 (1H, d,  $\downarrow$  8.0Hz), 7.56 (2H, app.d.  $\downarrow$  8.0Hz), 7.45 (1H, app.t,  $\downarrow$  8.0Hz), 7.21 (2H, d,  $\downarrow$  8.5Hz), 6.95 (2H, d,  $\downarrow$  8.5Hz), 5.17 (2H, s), 4.61-4.52 (1H, m), 3.11 (1H, dd,  $\downarrow$  13.8, 4.3Hz), 2.86 (1H, dd,  $\downarrow$  13.8, 10.4Hz), 2.31 (3H, s), 2.08 (3H, s), 2.07 (3H, s);  $m/z$  (ESI, 60V) 503 and 505 (MH<sup>+</sup>).

15 d) **O-(2,6-dichlorobenzyl)-N-(1-methyl-2-indolyl)-L-tyrosine**

from O-(2,6-dichlorobenzyl)-L-tyrosine methyl ester hydrochloride and 1-methylindole-2-carboxylic acid. Freeze drying afforded the title compound as a white amorphous solid (220mg).  $\delta$ H (DMSO-d<sub>6</sub>) 11.7 (1H, br s), 8.66 (1H, d,  $\downarrow$  8.3Hz), 7.65 (1H, d,  $\downarrow$  7.9Hz), 7.53 (2H, app.d,  $\downarrow$  8.1Hz), 7.26 (1H, obscured m), 7.09 (2H, app.t,  $\downarrow$  7.5Hz), 6.96 (2H, d,  $\downarrow$  8.5Hz), 5.16 (2H, s), 4.64-4.54 (1H, m), 3.89 (3H, s), 3.16 (1H, dd,  $\downarrow$  13.8, 4.3Hz) and 3.00 (1H, dd,  $\downarrow$  13.8, 10.4Hz);  $m/z$  (ESI, 60V) 497 and 499 (MH<sup>+</sup>).

25 e) **O-(2,6-dichlorobenzyl)-N-[2-(4-chlorophenyl)-3-(trifluoromethyl)-4-pyrazoyl]-L-tyrosine**

from O-(2,6-dichlorobenzyl)-L-tyrosine methyl ester hydrochloride and 2-(4-chlorophenyl)-3-(trifluoromethyl)-pyrazole-4-carboxylic acid. The title compound was isolated as an off-white solid (220mg).  $\delta$ H (DMSO-d<sub>6</sub>) 11.17 (1H, br s), 8.82 (1H, d,  $\downarrow$  8.2Hz), 8.08 (1H, s), 7.65 (2H, d,  $\downarrow$  8.1Hz), 7.54 (4H, app. d,  $\downarrow$  8.1Hz), 7.45 (1H, app. t,  $\downarrow$  8.0Hz), 7.24 (2H, d,  $\downarrow$  8.3Hz), 6.98 (2H, d,  $\downarrow$  8.3Hz), 5.18 (2H, s), 4.61-4.51 (1H, m), 3.17 (1H, dd,  $\downarrow$  13.8, 4.5Hz) and 2.94 (1H, dd,  $\downarrow$  13.8, 9.0Hz);  $m/z$  (ESI, 60V) 527 and 529 (MH<sup>+</sup>).

35 f) **O-(2,6-dichlorobenzyl)-N-(2-phenyl-4-thiazoyl)-L-tyrosine**



from O-(2,6-dichlorobenzyl)-L-tyrosine methyl ester hydrochloride and 2-phenyl-thiazole-4-carboxylic acid. Freeze-drying afforded the title compound as a pale yellow amorphous solid (340mg).  $\delta$ H (DMSO- $d_6$ ) 11.2 (1H, br s), 8.40 (1H, d,  $\downarrow$  8.1Hz), 8.31 (1H, s) 8.05-8.02 (2H, m), 7.55-7.51 (5H, m), 7.46-7.40 (1H, m), 7.23 (2H, d,  $\downarrow$  8.0Hz), 6.97 (2H, d,  $\downarrow$  7.8Hz), 5.16 (2H, s), 4.71-4.64 (1H, m), 3.18 (2H, app. d,  $\downarrow$  6.6Hz);  $m/z$  (ESI, 60V) 527 and 529 (MH<sup>+</sup>).

g) (N'-1-Methyl-5-nitropyrazolyl)-(N-2,6-dichlorobenzoyl)-L-4-aminophenylalanine

from (N-2,6-dichlorobenzoyl)-L-4-aminophenylalanine methyl ester with N-methyl-5-nitropyrazole-4-carboxylic acid.  $\delta$ H (DMSO- $d_6$ ) 10.64-(1H, s), 8.68 (1H, br d), 7.80 (1H, s), 7.64-7.48 (6H, m), 7.19 (2H, d,  $\downarrow$  8.5Hz), 4.59-4.49 (1H, m), 4.05 (3H, s), 3.18-2.9 (2H, m);  $m/z$  (ESI, 60V) 506 (MH<sup>+</sup>).

h) N-(2-Methylnicotinoyl)-O-(2,6-dichlorobenzyl) L-tyrosine

from O-(2,6-dichlorobenzyl)-L-tyrosine methyl ester hydrochloride and 2-methylnicotinic acid.  $\delta$ H (DMSO- $d_6$ ) 8.73 (1H, br d), 8.48 (1H, m), 7.7-7.46 (4H, m), 7.4-7.23 (3H, m), 7.00 (2H, d,  $\downarrow$  8.4Hz), 5.2 (2H, s), 4.60 (1H, m), 3.27-3.12 (1H, m), 3.0-2.82 (1H, m);  $m/z$  (ESI, 60V) 509 (MH<sup>+</sup>).

i) (N'-2-Chloronicotinoyl)-(N-benzoyl)-L-4-aminophenylalanine

from (N-benzoyl)-L-4-aminophenylalanine methyl ester and 2-chloronicotinic acid.  $\delta$ H (DMSO- $d_6$ ) 10.19 (1H, s), 8.96 (1H, d,  $\downarrow$  8.2Hz), 8.46 (1H, dd,  $\downarrow$  4.8, 1.9Hz), 7.96 (2H, dd,  $\downarrow$  6.7, 1.7Hz), 7.75-7.65 (3H, m), 7.62-7.45 (4H, m), 7.25 (2H, d,  $\downarrow$  8.5Hz), 4.6 (1H, m), 3.2-3.12 (1H, m), 3.0-2.89 (1H, m),  $m/z$  (ESI, 60V) 424 (MH<sup>+</sup>).

j) N'-(Quinoline-4-carbonyl)-O-(2,6-dichlorobenzyl)-L-tyrosine

from O-(2,6-dichlorobenzyl)-L-tyrosine methyl ester hydrochloride and 4-quinoline carboxylic acid.  $\delta$ H (DMSO- $d_6$ ) 9.0 (1H, d), 8.94 (1H, d,  $\downarrow$  4.3), 8.06 (1H, d,  $\downarrow$  8.2Hz), 7.79 (2H, m), 7.62-7.42 (4H, m), 7.34 (1H, d,  $\downarrow$  4.3Hz), 7.34 (2H, d,  $\downarrow$  8.7Hz), 7.00 (2H, d,  $\downarrow$  8.7Hz), 5.22 (2H, s), 4.72 (1H, m), 3.29-3.19 (1H, m), 3.0-2.88 (1H, m);  $m/z$  (ESI, 60V) 495 (MH<sup>+</sup>).

- k) *N'*-(2-Phenoxynicotinoyl)-O-(2,6-dichlorobenzyl)-L-tyrosine  
 from O-(2,6-dichlorobenzyl)-L-tyrosine methyl ester hydrochloride and 2-phenoxynicotinic acid.  $\delta$ H (DMSO- $d_6$ ) 8.55 (1H, d), 8.18 (2H, m), 7.6-7.4 (5H, m), 7.3-7.15 (6H, m), 6.71 (2H, d), 5.1 (2H, s), 4.65 (1H, m), 3.22-3.0 (2H, m);  $m/z$  (ESI, 60V) 537 (MH<sup>+</sup>).
- l) *N*-(Pyridine-2-carbonyl)-O-(2,6-dichlorobenzyl)-L-tyrosine  
 from O-(2,6-dichlorobenzyl)-L-tyrosine methyl ester hydrochloride and 2-picolinic acid.  $\delta$ H (DMSO- $d_6$ ) 8.8-8.6 (2H, m), 8.0 (2H, m), 7.7-7.4 (4H, m), 7.14 (2H, d,  $\downarrow$  8.7Hz), 6.92 (2H, d,  $\downarrow$  8.7Hz), 5.15 (2H, s), 4.72 (1H, m), 3.17 (2H, m);  $m/z$  (ESI, 60V) 445 (MH<sup>+</sup>).
- m) *N*-(Pyridine-4-carbonyl)-O-(2,6-dichlorobenzyl)-L-tyrosine  
 from O-(2,6-dichlorobenzyl)-L-tyrosine methyl ester hydrochloride and isonicotinic acid.  $\delta$ H (DMSO- $d_6$ ) 8.68 (2H, dd,  $\downarrow$  4.5, 1.6Hz), 8.4 (1H, d,  $\downarrow$  7.2Hz), 7.63 (2H, dd,  $\downarrow$  4.5, 1.6Hz), 7.6-7.4 (3H, m), 7.15 (2H, d,  $\downarrow$  8.6Hz), 6.87 (2H, d,  $\downarrow$  8.6Hz), 5.14 (2H, s), 4.31 (1H, m), 3.2-2.96 (2H, m);  $m/z$  (ESI, 60V) 445 (MH<sup>+</sup>).
- n) *N*-(2-Hydroxynicotinoyl)-O-(2,6-dichlorobenzyl)-L-tyrosine  
 from O-(2,6-dichlorobenzyl)-L-tyrosine methyl ester hydrochloride and 2-hydroxynicotinic acid.  $\delta$ H (DMSO- $d_6$ ) 10.15 (1H, d,  $\downarrow$  7.4Hz), 8.31 (1H, dd,  $\downarrow$  7.1, 2.0Hz), 7.71 (1H, br s), 7.65-7.45 (3H, m), 7.16 (2H, d,  $\downarrow$  8.4Hz), 6.97 (2H, d,  $\downarrow$  8.4Hz), 6.46 (1H, t,  $\downarrow$  6.8Hz), 5.17 (2H, s), 4.68 (1H, m), 3.2-3.0 (2H, m);  $m/z$  (ESI, 60V) 461 (MH<sup>+</sup>).
- o) *(N'*-(2-Aminonicotinoyl)-(*N*-2,6-dichlorobenzoyl)-L-4-aminophenylalanine  
 from (*N*-2,6-dichlorobenzoyl)-L-4-aminophenylalanine methyl ester and 2-aminonicotinic acid.  $\delta$ H (DMSO- $d_6$ ) 10.64 (1H, s), 8.55 (1H, d), 8.06 (1H, m), 7.8 (1H, d), 7.7-7.4 (4H, m), 7.28 (2H, d,  $\downarrow$  8.5Hz), 6.9 (2H, br s), 6.5 (1H, m), 4.52 (1H, m), 3.2-2.9 (2H, m);  $m/z$  (ESI, 60V) 473 (MH<sup>+</sup>).
- p) *(N'*-(2-Hydroxynicotinoyl)-(*N*-3,5-dichloroisonicotinoyl)-L-4-aminophenylalanine

from (N-3,5-dichloroisonicotinoyl)-L-4-aminophenylalanine methyl ester and 2-hydroxynicotinic acid.  $\delta$ H (DMSO- $d_6$ ) 10.14 (1H, d,  $\downarrow$  7.6Hz), 8.78 (2H, s), 8.31 (1H, dd,  $\downarrow$  7.2, 2.2Hz), 7.70 (1H, br m), 7.57 (2H, d,  $\downarrow$  8.5Hz), 7.20 (2H, d,  $\downarrow$  8.5Hz), 6.46 (1H, m), 4.69 (1H, m), 3.20-2.95 (2H, m);  $m/z$  (ESI, 60V) 475 (MH<sup>+</sup>).

q) (N'-2-Methylnicotinoyl)-(N-3,5-dichloroisonicotinoyl)-L-4-aminophenylalanine

from (N-3,5-dichloroisonicotinoyl)-L-4-aminophenylalanine and 2-methylnicotinic acid.  $\delta$ H (DMSO- $d_6$ ) 8.78 (2H, s), 8.60 (1H, br d), 8.48 (1H, dd,  $\downarrow$  4.8, 1.6Hz), 7.7-7.5 (3H, m), 7.4-7.2 (3H, m), 4.5 (1H, ), 3.3-2.85 (2H, m), 2.34 (3H, s);  $m/z$  (ESI, 60V) 473 (MH<sup>+</sup>).

r) (N'-2,6-Dichlorobenzoyl)-(N-2-phenoxy nicotinoyl)-L-4-aminophenylalanine

from (N-2,6-dichlorobenzoyl)-L-4-aminophenylalanine methyl ester and 2-phenoxy nicotinic acid.  $\delta$ H (DMSO- $d_6$ ) 10.61 (1H, s), 8.56 (1H, d,  $\downarrow$  7.5Hz), 8.22-8.17 (2H, m), 7.59-7.39 (8H, m), 7.27-7.11 (7H, m), 4.72-4.64 (1H, m, CH), 3.16 (1H, dd,  $\downarrow$  13.7, 4.9Hz, CH<sub>A</sub>H<sub>B</sub>) and 3.06 (1H, dd,  $\downarrow$  13.7, 7.8Hz, CH<sub>A</sub>H<sub>B</sub>);  $m/z$  (ESI, 60V) 550 (MH<sup>+</sup>).

s) (N-3,5-Dichloroisonicotinoyl)-(N-2-phenoxy nicotinoyl)-L-4-aminophenylalanine

from (N-3,5-dichloroisonicotinoyl)-L-4-aminophenylalanine methyl ester and 2-phenoxy nicotinic acid.  $\delta$ H (DMSO- $d_6$ ) 10.79 (1H, s), 8.78 (2H, s, ArH), 8.57 (1H, d,  $\downarrow$  7.6Hz), 8.25-8.13 (2H, m), 7.46-7.35 (4H, m), 7.29-7.05 (6H, m), 4.73-4.62 (1H, m, CH), 3.17 (1H, dd,  $\downarrow$  13.7, 4.7Hz, CH<sub>A</sub>H<sub>B</sub>) and 3.09 (1H, dd,  $\downarrow$  13.7, 7.8Hz, CH<sub>A</sub>H<sub>B</sub>);  $m/z$  (ESI, 60V) 551 (MH<sup>+</sup>).

t) N-(4-Acetyl-1,2,5-trimethyl-1H-pyrrole-3-carbonyl)-N-(3,5-dichloro-4-picolyl)-L-4-aminophenylalanine

from (N-3,5-dichloro-4-picolyl)-L-4-aminophenylalanine methyl ester and 4-acetyl-1,2,5-trimethyl-1H-pyrrole-3-carboxylic acid.  $\delta$ H (DMSO- $d_6$ , 300MHz) 400K) 12.72 (1H, br s), 10.83 (1H, s), 8.80 (2H, s), 8.42 (1H, d,  $\downarrow$  8.4Hz), 7.54 (2H, d,  $\downarrow$  8.5Hz), 7.30 (2H, d,  $\downarrow$  8.4Hz), 4.66 (1H, m), 3.34

(3H, s), 3.16 (1H, dd,  $\downarrow$  13.8, 4.3Hz), 2.89 (1H, dd,  $\downarrow$  13.8, 11.0Hz), 2.32 (3H, s), 2.05 (3H, s) and 1.90 (3H, s);  $m/z$  (ESI, 60V) 531 (MH<sup>+</sup>).

5 **u) N-(4-(Carboxy)nicotinoyl)-N'-(3,5-dichloro-4-picolyl)-L-4-aminophenylalanine**

from (N-3,5-dichloro-4-picolyl)-L-4-aminophenylalanine methyl ester and 4-(methoxycarbonyl)nicotinic acid.  $\delta$ H (DMSO-d<sub>6</sub>, 300MHz) 10.98 (1H, s), 9.01 (1H, d,  $\downarrow$  8.0Hz), 8.77 (2H, s), 8.63 (1H, s), 7.66 (1H, d,  $\downarrow$  5.1Hz), 7.59 (2H, d,  $\downarrow$  8.5Hz), 7.32 (2H, d,  $\downarrow$  8.5Hz), 4.63 (1H, m), 3.14 (1H, dd,  $\downarrow$  13.9, 5.4Hz) and 3.03 (1H, dd,  $\downarrow$  13.9, 8.8Hz);  $m/z$  (ESI, 160V) 503 (MH<sup>+</sup>).

15 **v) (2-Acetyl-3-thienyl)carbonyl-(N-3,5-dichloro-4-picolyl)-L-4-aminophenylalanine**

from (N-3,5-dichloro-4-picolyl)-L-4-aminophenylalanine methyl ester and 2-acetyl-thiophene-3-carboxylic acid.  $\delta$ H (DMSO-d<sub>6</sub>) 9.17 (1H, d,  $\downarrow$  8.2Hz, NH), 8.77 (2H, s, pyr-H), 7.99 (1H, d,  $\downarrow$  5.1Hz, thiophene H-5), 7.57 (2H, ABd,  $\downarrow$  8.5Hz, Ar-H), 7.31 (2H, ABd, 2H,  $\downarrow$  8.5Hz, Ar-H), 7.07 (1H, d,  $\downarrow$  5.1Hz, thiophene H-4), 5.38 (1H, m, CH $\alpha$ ) 3.18 (1H, dd,  $\downarrow$  13.8, 4.6Hz, CHCH<sub>A</sub>H<sub>B</sub>Ar), 2.93 (1H, dd,  $\downarrow$  13.8, 10.3Hz, CHCH<sub>A</sub>H<sub>B</sub>Ar) and 2.26 (3H, s, COMe);  $m/z$  (ESI, 60V) 506 (MH<sup>+</sup>).

**EXAMPLE 6**

25 **a) 2-Chloronicotinoyl-(N-2,6-dichlorobenzoyl)-L-4-aminophenylalanine**

Lithium hydroxide monohydrate (109mg, 2.5mmol) was added to a solution of Intermediate 8 (450mg, 1mmol) in a mixture of THF (10ml) and water (10ml). The mixture was stirred for 2h at room temperature, then the THF was evaporated *in vacuo*. The aqueous residue was neutralised (1M hydrochloric acid), and the precipitate isolated by filtration, washed with water and dried to give the title compound (300mg, 61%).  $\delta$ H (DMSO-d<sub>6</sub>, 400K) 8.90 (1H, d,  $\downarrow$  9.1Hzm 1NH), 8.48 (1H, m, Py H), 7.69 (1H, m, Py H), 7.64-7.45 (7H, m, 4ArH, 1 NH, 1 PyH), 7.29 (2H, d,  $\downarrow$  8.3Hz, 2 ArH), 4.66-4.53 (1H, m, CH $\alpha$ tyr), 3.18 (1H, dd,  $\downarrow$  14.52Hz CH<sub>A</sub>H<sub>B</sub>Ar), and 2.91 (1H, dd,  $\downarrow$  9.6, 14Hz, CH<sub>A</sub>H<sub>B</sub>Ar).  $m/z$  (ESI, 60V) 491 (MH<sup>+</sup>).

35

The following compounds were prepared in a similar manner:

b) **N-(2-Chloronicotinoyl)-O-(2,6-dichlorobenzoyl)-L-tyrosine**

from Intermediate 30 to give the title compound as a white solid.  $\delta$ H (DMSO- $d_6$ , 300MHz) 12.91 (1H, br s, CO<sub>2</sub>H), 8.98 (1H, d,  $J$  8.3Hz, CONH), 8.44 (1H, dd,  $J$  4.8, 1.9Hz, PyH), 7.69-7.58 (4H, m, Cl<sub>2</sub>ArH<sub>3</sub> + PyH), 7.45 (1H, dd,  $J$  7.5, 4.9Hz, PyH), 7.43 (2H, d,  $J$  8.7Hz, ArH), 7.22 (2H, d,  $J$  8.5Hz, ArH), 4.67 (1H, ddd,  $J$  10.0, 8.2, 4.7Hz, CH $\alpha$ ), 3.24 (1H, dd,  $J$  14.0, 4.0Hz, CH<sub>A</sub>H<sub>B</sub>Ar) and 2.99 (1H, dd,  $J$  13.9, 10.2Hz, CH<sub>A</sub>H<sub>B</sub>Ar);  $m/z$  (ESI, 60V) 493 (MH<sup>+</sup>).

c) **N-(2-Chloronicotinoyl)-N-methyl-N'-(3,5-dichloro-4-picolyl)-L-4-aminophenylalanine**

from Intermediate 31 to give the title compound as a white solid  $\delta$ H (DMSO- $d_6$ , 300MHz, 405K) 10.36 (1H, br s, CONH), 8.67 (2H, s, Cl<sub>2</sub>PyH), 8.42-8.39 (1H, m, ClPyH), 7.54-7.15 (6H, br m, 4 x ArH + 2 x ClPyH), 5.30 (1H, v br s, CH $\alpha$ ), 3.4-2.6 (5H, br m, NMe + CHCH<sub>2</sub>Ar) (Acid proton not observed at 405K, at 300K  $\delta$ H 13.06 (1H, br s, CO<sub>2</sub>H));  $m/z$  (ESI, 70V) 507 (MH<sup>+</sup>).

d) **[(S-2,5-dimethoxyphenyl)sulphonyl]nicotinoyl-O-(2,6-dichlorobenzyl)-L-tyrosine**

from Intermediate 32  $\delta$ H (DMSO- $d_6$ ) 8.57 (2H, m, pyrH, NH), 7.83 (1H, d,  $J$  7.8Hz, pyr-H), 7.69 (1H, dd,  $J$  7.8, 4.7Hz, pyr-H), 7.55 (4H, m, Ar-H), 7.28-7.22 (3H, m, Ar-H), 7.07 (1H, d,  $J$  9.1Hz, Ar-H), 6.95 (2H, d,  $J$  8.5Hz, Ar-H), 5.18 (1H, s, CH<sub>2</sub>O), 4.55 (1H, m, CH $\alpha$ ), 3.81 (3H, s, OMe), 3.40 (3H, s, OMe) and 3.07 (2H, m, CHCH<sub>2</sub>Ar).  $m/z$  (ESI, 30V) 645 (MH<sup>+</sup>).

e) **2-[(2-Chloro-3-pyridinyl)carbonyl]amino-3-(4-[(2,6-dichloroanilino)carbonyl]phenyl)propanoic acid**

from Intermediate 15 to give the title compound as an off white solid  $\delta$ H (DMSO- $d_6$ , 300K) 11.06 (1H, br s), 10.23 (1H, br s), 9.00 (1H, d,  $J$  7.8Hz), 8.46 (1H, br d), 7.95 (2H, d,  $J$  7.4Hz), 7.69 (1H, d,  $J$  7.1Hz), 7.59 (2H, d,  $J$  8.0Hz), 7.50-7.36 (3H, m), 4.71 (1H, br), 3.26 (1H) and 3.06 (1H, dd,  $J$  13.8, 10.0Hz);  $m/z$  (ESI, 60V) 492 (MH<sup>+</sup>).

f) **2-(((2-Chloro-3-pyridinyl)carbonyl)amino)-3-(4-(((3,5-dichloro-4-pyridinyl)amino)carbonyl))phenyl propionic acid**

from the corresponding intermediate ester prepared in a similar way to Intermediate 15 to give title compounds as an offwhite solid.  $\delta$ H (DMSO- $d_6$ , 300MHz) 12.92 (1H, br s), 10.57 (1H, s), 9.00 (1H, d,  $\downarrow$  8.2Hz), 8.75 (2H, s), 8.45 (1H, dd,  $\downarrow$  4.7, 1.7Hz), 7.96 (2H, d,  $\downarrow$  8.15Hz), 7.69 (1H, d,  $\downarrow$  7.3, 1.7Hz), 7.49 (2H, d,  $\downarrow$  8.0Hz), 7.48 (1H, 4.71 (1H, br m), 3.28 (1H, dd,  $\downarrow$  13.9, 4.7Hz) and 3.06 (1H, dd,  $\downarrow$  13.9, 10.0Hz);  $m/z$  (ESI, 60V) 483 (MH<sup>+</sup>).

**EXAMPLE 7**

**2-Thio(S-acetic acid)nicotinoyl-O-(2,6-dichlorobenzyl)-L-tyrosine**

Lithium hydroxide monohydrate (75mg, 1.8mmol) was added to a solution of Intermediate 9 (360mg, 0.6mmol) in a mixture of THF (13ml) and water (10ml). The mixture was stirred for 2hr at room temperature, then the THF was evaporated *in vacuo*. The aqueous residue was neutralised (1M hydrochloric acid), and the precipitate isolated by filtration, washed with water and dried to give the title compound (200mg, 58%).  $\delta$ H (DMSO- $d_6$ , 400K), 8.5-8.35 (2H, m, pyrH, 1NH), 7.71 (1H, dd,  $\downarrow$  1.7, 7.6, pyrH), 7.57 (2H, d,  $\downarrow$  8.9, 2ArH), 7.45 (1H, m, 1ArH), 7.22 (2H, d,  $\downarrow$  8.5, 2ArH), 7.18 (1H, m, 1pyrH), 6.97 (2H, d,  $\downarrow$  8.5, 2ArH), 5.18 (2H, s, OCH<sub>2</sub>Ar), 4.43 (1H, m, CH<sub>2</sub>tyr), 3.82 (2H, s, SCH<sub>2</sub>CO<sub>2</sub>H), 2.94-3.23 (2H, m, CH<sub>2</sub>Ar);  $m/z$  (ESI, GOU) 535 (MH<sup>+</sup>).

**EXAMPLE 8**

**2-Thio(S-methyl)nicotinoyl-O-(2,6-dichlorobenzyl)-L-tyrosine**

Lithium hydroxide monohydrate (140mg, 3.3mmol) was added to a solution of Intermediate 10 (1.4gm, 2.7mmol) in a mixture of THF (10ml) and water (10ml). The mixture was stirred for 2 hr at room temperature then the THF was evaporated *in vacuo*. The aqueous residue was neutralised (1M hydrochloric acid), and the precipitate isolated by filtration, washed with water and dried to give the title compound (1.1gm, 81%).  $\delta$ H (DMSO- $d_6$ , 400K) 8.73 (1H, d,  $\downarrow$  8.1, NH), 8.52 (1H, dd,  $\downarrow$  1.7, 4.8, 1pyrH), 7.67 (1H, dd,  $\downarrow$  1.7, 7.6, 1pyrH), 7.55 (2H, d,  $\downarrow$  8.9, 2ArH), 7.45 (1H, dd,  $\downarrow$  2.3, 8.9, 1ArH), 7.24 (2H, d,  $\downarrow$  8.6, 2ArH), 7.16 (1H, m, 1pyrH), 6.99 (2H, d,  $\downarrow$  8.6,

2ArH), 5.18 (2H, s, OCH<sub>2</sub>Ar), 4.55 (1H, m, CH<sub>2</sub>tyr), 3.16-2.95 (2H, m, CH<sub>2</sub>Ar) and 2.38 (3H, s, SCH<sub>3</sub>); *m/z* (ESI, GOU) 491 (MH<sup>+</sup>).

#### **EXAMPLE 9**

5 **(N-2-Chloronicotinoyl)-4-[(E)-2-(2,6-dichlorophenyl)ethenyl]**  
**phenylalanine**

To a solution of Intermediate 28 (1.0g, 1.99mmol) in THF (5ml) and water (5ml) was added lithium hydroxide monohydrate (88mg 2.09mmol). The reaction mixture was stirred for 1h. The THF was then removed *in vacuo* and the remaining aqueous solution acidified to pH6 with 1M hydrochloric acid. The resulting precipitate was collected and washed with water and ether and finally freeze dried. The resulting compound contained an impurity so a small amount was purified by preparative HPLC (98mg).  $\delta$ H (DMSO-d<sub>6</sub>) 12.90 (1H, dr s, CO<sub>2</sub>H), 8.98 (1H, d,  $\downarrow$  8.0Hz), 8.45 (1H, d,  $\downarrow$  3.4Hz), 7.70-7.03 (9H, m), 4.72-4.60 (1H, m, CH), 3.20 (1H, dd,  $\downarrow$  14.0, 4.5Hz, CH<sub>A</sub>H<sub>B</sub>) and 3.01 (1H, dd,  $\downarrow$  14.0, 9.9Hz, CH<sub>A</sub>H<sub>B</sub>); *m/z* (ESI, 60V) 475 (MH<sup>+</sup>).

#### **EXAMPLE 10**

20 **(N-2-Chloronicotinoyl)-4-[2-(2,6-dichlorophenyl)-2-hydroxyethyl]**  
**phenylalanine**

Lithium hydroxide monohydrate (13mg) was added to a solution of Intermediate 22 (150mg, 0.29mmol) in THF (5ml) and H<sub>2</sub>O (5ml). The solution was stirred for 1h and then the THF removed *in vacuo* and the remaining aqueous solution acidified to pH6 with 1M hydrochloric acid. The solid precipitate formed was collected by filtration, washed with copious quantities of water and finally freeze dried to give the title compound as a fluffy white solid (70mg, 49%).  $\delta$ H (DMSO-d<sub>6</sub>) 12.78 (1H, br s, CO<sub>2</sub>H), 8.90 (1H, d,  $\downarrow$  8.0Hz), 8.45 (1H, d,  $\downarrow$  4.8Hz), 7.63-7.58 (1H, m, ArH), 7.49-7.45 (1H, m, ArH), 7.34-7.07 (7H, m, ArH), 5.50-5.45 (1H, m, CH), 4.61-4.51 (1H, m, CH) and 3.30-2.35 (4H, m, 2 x CH<sub>2</sub>); *m/z* (ESI, 60V) 493 (MH<sup>+</sup>).

#### **EXAMPLE 11**

35 **(N-2-Chloronicotinoyl)-4-[2-(2-dichlorophenyl)-2-oxoethyl]**  
**phenylalanine**

Lithium hydroxide monohydrate (36mg, 0.85mmol) was added to a solution of Intermediate 23 (400mg, 0.77mmol) in THF (5ml) and water (5ml). The reaction mixture was stirred for 3h and then the THF was removed *invacuo*. The remaining aqueous solution was acidified with 1M hydrochloric acid. The resulting white precipitate was collected and washed well with water. Further purification by column chromatography (SiO<sub>2</sub>; acetic acid: MeOH:DCM, 2:8:90) gave the title compound as a white solid (78mg, 19%).  $\delta$ H (DMSO-d<sub>6</sub>) 8.45 (1H, dd,  $\downarrow$  4.8, 2.0Hz), 8.03 (1H, dd,  $\downarrow$  7.7, 2.0Hz), 7.36-7.20 (8H, m, ArH), 6.98 (1H, d,  $\downarrow$  7.2Hz, ArH), 5.14-5.05 (1H, m, CH), 4.11 (2H, s, CH<sub>2</sub>C=O)m, 3.39 (1H, dd,  $\downarrow$  14.1, 5.6Hz, CH<sub>A</sub>H<sub>B</sub>) and 3.25 (1H, dd,  $\downarrow$  14.1, 6.2Hz, CH<sub>A</sub>H<sub>B</sub>);  $m/z$  (ESI, 60V) 491 (MH<sup>+</sup>).

#### $\alpha_4\beta_1$ Integrin-dependent Jurkat cell adhesion to VCAM-Ig

96 well NUNC plates were coated with F(ab)<sub>2</sub> fragment goat anti-human IgG Fc $\gamma$ -specific antibody [Jackson Immuno Research 109-006-098: 100  $\mu$ l at 2  $\mu$ g/ml in 0.1M NaHCO<sub>3</sub>, pH 8.4], overnight at 4°. The plates were washed (3x) in phosphate-buffered saline (PBS) and then blocked for 1h in PBS/1% BSA at room temperature on a rocking platform. After washing (3x in PBS) 9 ng/ml of purified 2d VCAM-Ig diluted in PBS/1% BSA was added and the plates left for 60 minutes at room temperature on a rocking platform. The plates were washed (3x in PBS) and the assay then performed at 37° for 30 min in a total volume of 200  $\mu$ l containing 2.5 x 10<sup>5</sup> Jurkat cells in the presence or absence of titrated test compounds.

Each plate was washed (2x) with medium and the adherent cells were fixed with 100 $\mu$ l methanol for 10 minutes followed by another wash. 100 $\mu$ l 0.25% Rose Bengal (Sigma R4507) in PBS was added for 5 minutes at room temperature and the plates washed (3x) in PBS. 100 $\mu$ l 50% (v/v) ethanol in PBS was added and the plates left for 60min after which the absorbance (570nm) was measured.

#### $\alpha_4\beta_7$ Integrin-dependent JY cell adhesion to MAdCAM-Ig

This assay was performed in the same manner as the  $\alpha_4\beta_1$  assay except that MAdCAM-Ig (150ng/ml) was used in place of 2d VCAM-Ig and a sub-line of the  $\beta$ -lympho blastoid cell-line JY was used in place of Jurkat cells.



The IC<sub>50</sub> value for each test compound was determined as described in the  $\alpha_4\beta_1$  integrin assay.

**$\alpha_5\beta_1$  Integrin-dependent K562 cell adhesion to fibronectin**

- 5 96 well tissue culture plates were coated with human plasma fibronectin (Sigma F0895) at 5 $\mu$ g/ml in phosphate-buffered saline (PBS) for 2 hr at 37°C. The plates were washed (3x in PBS) and then blocked for 1h in 100 $\mu$ l PBS/1% BSA at room temperature on a rocking platform. The blocked plates were washed (3x in PBS) and the assay then performed at 10 37°C in a total volume of 200 $\mu$ l containing 2.5x 10<sup>5</sup> K562 cells, phorbol-12-myristate-13-acetate at 10ng/ml, and in the presence or absence of titrated test compounds. Incubation time was 30 minutes. Each plate was fixed and stained as described in the  $\alpha_4\beta_1$  assay above.

15  **$\alpha_m\beta_2$ -dependent human polymorphonuclear neutrophils adhesion to plastic**

- 96 well tissue culture plates were coated with RPMI 1640/10% FCS for 2h at 37°C. 2 x 10<sup>5</sup> freshly isolated human venous polymorphonuclear neutrophils (PMN) were added to the wells in a total volume of 200 $\mu$ l in the presence of 10ng/ml phorbol-12-myristate-13-acetate, and in the presence or absence of test compounds, and incubated for 20min at 37°C followed by 30min at room temperature. The plates were washed in medium and 100 $\mu$ l 0.1% (w/v) HMB (hexadecyl trimethyl ammonium bromide, Sigma H5882) in 0.05M potassium phosphate buffer, pH 6.0 added to each well. 20 25 The plates were then left on a rocker at room temperature for 60 min. Endogenous peroxidase activity was then assessed using tetramethyl benzidine (TMB) as follows: PMN lysate samples mixed with 0.22% H<sub>2</sub>O<sub>2</sub> (Sigma) and 50 $\mu$ g/ml TMB (Boehringer Mannheim) in 0.1M sodium acetate/citrate buffer, pH 6.0 and absorbance measured at 630nm.

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**$\alpha_{IIb}\beta_3$ -dependent human platelet aggregation**

- Human platelet aggregation was assessed using impedance aggregation on the Chronolog Whole Blood Lumiaggregometer. Human platelet-rich plasma (PRP) was obtained by spinning fresh human venous blood anticoagulated with 0.38% (v/v) tri-sodium citrate at 220xg for 10 min and diluted to a cell density of 6 x 10<sup>8</sup>/ml in autologous plasma. Cuvettes 35

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